

Wed Nov 7 09:21:19 2001

us-09-656-668-198.ali.rst

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: November 5, 2001, 22:35:55 ; Search time 1157.88 Seconds
(without alignments)
4947.343 Million cell updates/sec

Title: US-09-656-668-198
Perfect score: 606
Sequence: 1 tgaagttgcccttaccctccccc.....aagcctgtcttctgtctgcac 606

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 10228115 seqs, 4736426750 residues

Word size : 0

Total number of hits satisfying chosen parameters: 20456230

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

EST.*

1: gb_est1.*

2: gb_est2.*

3: gb_est3.*

4: gb_est4.*

5: gb_est5.*

6: gb_est6.*

7: gb_est7.*

8: gb_est8.*

9: gb_est9.*

10: gb_est10.*

11: gb_est11.*

12: gb_est12.*

13: gb_est13.*

14: gb_est14.*

15: gb_est15.*

16: gb_est16.*

17: gb_est17.*

18: gb_est18.*

19: gb_est19.*

20: gb_est20.*

21: gb_est21.*

22: gb_est22.*

23: gb_est23.*

24: gb_est24.*

25: gb_est25.*

26: gb_est26.*

27: gb_est27.*

28: gb_est28.*

29: gb_est29.*

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31: gb_est31.*

32: gb_est32.*

33: gb_est33.*

34: gb_est34.*

35: gb_est35.*

36: gb_est36.*

37: gb_est37.*

38: gb_est38.*

39: gb_est39.*

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41: gb_est41.*

42: gb_est42.*

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44: gb_est44.*

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46: gb_est46.*

47: gb_est47.*

48: gb_est48.*

49: gb_est49.*

50: gb_est50.*

44: em_esthum10.*

45: em_esthum11.*

46: em_esthum12.*

47: em_esthum13.*

48: em_esthum14.*

49: em_esthum15.*

50: em_esthum16.*

51: em_esthum17.*

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53: em_esthum19.*

54: em_esthum20.*

55: em_esthum21.*

56: em_esthum22.*

57: em_esthum23.*

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59: em_esthum25.*

60: em_esthum26.*

61: em_esthum27.*

62: em_esthum28.*

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66: em_esthum32.*

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101: em_esthum67.*

102: em_esthum68.*

103: em_esthum69.*

104: em_esthum70.*

105: em_esthum71.*

106: em_esthum72.*

107: em_esthum73.*

108: em_esthum74.*

109: em_esthum75.*

110: em_esthum76.*

111: em_esthum77.*

112: em_esthum78.*

113: em_esthum79.*

114: em_esthum80.*

115: em_esthum81.*

116: em_esthum82.*

117: gb_est48:*
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193: em_gss_fun:*
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255: em_gss_inv53:*
256: em_gss_inv54:*
257: em_gss_inv55:*
258: em_gss_inv56:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

Unpublished (1997)
Contact: Robert Strausberg, Ph.D.

source

BASE COUNT
ORIGIN

Query Match: 99.7%; Pred. No. 1.1e-158;
Best Local Similarity 0; Mismatches 1; Indels 0; Gaps 0;
Conservative 0

Matches	363;	Conservative	0;	Mismatches	1;	Insertions	0;	Deletions	0;
---------	------	--------------	----	------------	----	------------	----	-----------	----

408 TGAGAAACCTACCGCAGGATCTTACTGGCCTTCATAGGTAAGCTTGCCTTGTCTCGGC 342

TCTTTTCCATCAGGTCCTCAACGTCCCGAGCCAGG 285

[illegible]

380 aactagaagaatgaccaaccatcgtgtgacctggactgtccctagctllcagcaccgaa

...tctcgttcgaagccctcaaacctaaactatgacatacaccctagcagctgagg 49

DB

I00

E8

Db 108 ACTCTCAATACGAATTAGTCTTTGTCACCTGGAGATGAAATAAC... ..

18 mcm 15

RESULT 2

DEFINITION od71f05.s1 NCI_CGAP_OV2 Homo sapiens CDNA clone IMAGE:30000000

```

VERSION      AA62630/:1  01:202000
KEYWORDS     EST

```

Result	Query
No.	Score Match

45	19	3.1
44	19	3.1
c		

ALIGNMENTS

clone IMZ
repetitive

ACCESSION	AI023799
VERSION	AI023799

KEYWORDS EST.
SOURCE human.

ORGANISM	Homo sapiens	Eukaryotes
...

REFERENCE
AUTHORS
NCT-CCAP
Mammalia,
1 (bases

AUTHORS	NEL CGAF
TITLE	National

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SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 405)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-remail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Michael R. Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: David B. Kitzman, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LINL at: www.bio.lnlnl.gov/dbp/image/image.html
Insert length: 500 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham.
Location/Qualifiers
FEATURES
source
1. 405
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1373409"
/clone.lib="NCI-CGAP_OV2"
/sex="female"
/tissue.type="ovary"
/lab.host="DH10B"
/note="Vector: pAMP10; mRNA made from invasive ovarian tumor, cDNA made by oligo-dr priming. Non-directionally cloned. Size-selected on agarose gel, average insert size 600 bp. Reference: Krizman et al. (1996) Cancer Research 56:5380-5383."
BASE COUNT 118 a 102 c 79 g 106 t
ORIGIN

Query Match 45.4%; Score 275; DB 12; Length 405;
Best Local Similarity 99.7%; Pred. No. 4.5e-138;
Matches 325; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY 272 ataaataagaacatgcacagtcctccctccctaaagtcctccgagcgaagcgtcaaggaat 331
|||||
DB 9 ATAAATAAGACATGCGCCAGTCCCTCCCTCACGCTCCGACGAGCTTCAAGCAAT 68
OY 332 tccataacagtagaagaacactaaatattgattcaaaaatctcaagcaactagaagaat 391
|||||
DB 69 TCCAAATTAAGAGTAAATATGATTTCAAAAATCTCAGCACTGAGAGAAAT 128
OY 392 gacaaacactcctgtgtgctgctgagctgtcctagtttagcatggaagtttcaagttc 451
|||||
DB 129 GACCAACATCTCGTTGGCTGGAGCTGCTAGTTTACGATTGAAGTTTCAGTTTC 188
OY 452 caggaagcctcaagcctggtgctgtgtcaccctcagagctgagggactctcaatac 511
|||||
DB 189 CAGGAAGCCCTCAGGCTCGGCTCTGCTACCTGACCTGAGGAGCACTTCAATAC 248
OY 512 agaattgtcttctgtcactgtagaataatacttaatttgtaacatgtgaaacatc 571
|||||
DB 249 AGAATTGTCTTTGCCCTGAGATGAATTAATTGTTTACATGATGAAACATC 308
OY 572 tataaacctactgaagcctgtct 597
|||||
DB 309 TATTAACATCTACTGAAGCCTGTCT 334

RESULT 3
AI683094 473 bp mRNA EST 16-DEC-1999
LOCUS AI683094/c
DEFINITION tx01et12.x1 NCI-CGAP_Ut4 Homo sapiens cDNA clone IMAGE:2267950 3',
MRNA sequence

ACCESSION AI683094
VERSION AI683094.1 GI:4893286
KEYWORDS EST.
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 473)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-remail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LINL at: www.bio.lnlnl.gov/dbp/image/image.html
Insert length: 2912 Std Error: 0.00
Seq primer: -400P from Gibco
High quality sequence stop: 405.
Location/Qualifiers
FEATURES
source
1. 473
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2267950"
/clone.lib="NCI-CGAP_Ut4"
/tissue.type="serous papillary carcinoma, high grade, 2 pooled tumors"
/lab.host="DH10B"
/note="Organ: uterus; Vector: pGMV-SPORT6; Site:1; Salt: 0.5M; pH: 8.0; Cloned unidirectionally. Primer: Oligo dr. Site:2; NotI; Cloned unidirectionally. Primer: Oligo dr. Average insert size 1.48 kb. Life Technologies catalog #:
11542-016"
BASE COUNT 142 a 88 c 95 g 148 t
ORIGIN

Query Match 44.2%; Score 268; DB 23; Length 473;
Best Local Similarity 100.0%; Pred. No. 2.9e-134;
Matches 268; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 330 attccaatagtagaatagaacactaaatattgattcaaaaatctcaagcaactagaaga 389
|||||
DB 473 ATTCCAAATTAAGAGTAAATATGATTTCAAAAATCTCAGCACTGAGAGAA 414
OY 390 atgaccaacactcctgtgtgctgctgagctgtcctagtttagcatggaagtttcaagtt 449
|||||
DB 413 ATGACCAACCATCTCGTGTGCTGCTGAGCTGCTAGTTTACGATTGAAGTTTCAGT 354
OY 450 tccaggaagcctcagcctggtgctgtgtcaccctcagagctgagggactcttcaat 509
|||||
DB 353 TCCAGGAAGCCTCAGGCTCGGCTGCTGCTACCTGACGCTGAGGAGCACTTCAAT 294
OY 510 acaggaattgtctgtgctgctgagtagaataatacttaatttgtaacatgtgaaaca 569
|||||
DB 293 ACAGAAATTAAGCTTTGTGCTGCTGAGATGAATTAATTGTAACATGTAAGAAACA 234
OY 570 tctataaacactcactgaagcctgtct 597
|||||
DB 233 TCTATTAACATCTACTGAGCCTGTCT 206

RESULT 4
AI307373 414 bp mRNA EST 08-APR-1999
LOCUS AI307373/c
DEFINITION tb26f01.x1 NCI-CGAP_K1d12 Homo sapiens cDNA clone IMAGE:2055461 3',
MRNA sequence.
ACCESSION AI307373

VERSION	AI107373.1	GI:4002098
KEYWORDS	EST.	
SOURCE	human.	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.	
REFERENCE	1 (bases 1 to 414)	
FEATURES	NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap .	
FEATURES	National Cancer Institute, Cancer Genome Anatomy Project (CGAP),	
FEATURES	Tumor Gene Index	
COMMENT	Unpublished (1997)	
COMMENT	Contact: Robert Strausberg, Ph.D.	
COMMENT	Email: cgaps-r@nhi.nih.gov	
COMMENT	Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.	
COMMENT	Emmert-Buck, M.D., Ph.D.	
COMMENT	cDNA Library Preparation: M. Bento Soares, Ph.D.	
COMMENT	cDNA Library Arrayed by: Greg Lennon, Ph.D.	
COMMENT	DNA Sequencing by: Washington University Genome Sequencing Center	
COMMENT	Clone distribution: NCI-CGAP clone distribution information can be	
COMMENT	found through the I.M.A.G.E. Consortium/BLM at:	
COMMENT	www.bio.lnl.gov/bbrp/image/image.html	
COMMENT	Insert length: 592	
COMMENT	Std Error: 0.00	
COMMENT	Seq primer: -40UP from Gibco.	

```

FEATURES
SOURCE
1. .414
Location/Genbank accession number
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_image="2055481"
/clone_lib="NCI_CGAP_Kid12"
/tissue_type="2 pooled tumors (clear cell type)"
/lab_host="DH10B"
/note="Organ: Kidney; Vector: pF713D-Pac (Pharmacia) with
a modified polylinker; Site.1: Not I; Site.2: Eco RI;
Plasmid DNA from the normalized library NCI_CGAP_Kid5 was
prepared, and ss circles were made in vitro. Following HAP
purification, this DNA was used as tracer in a subtractive
hybridization reaction. The driver was PCR-amplified cDNAs
from a pool of 5,000 clones made from the same library
(cloneids 1333912-1325831, 1471368-1472903 and
1492104-1493555). Subtraction by Bento Soares and M.
Fatima Bernaldo."
BASE COUNT
126 a 80 c 85 g 123 t
ORIGIN

```

	Query Match	33.7%	Score 204;	DB 18:	Length 414;
	Best local Similarity	100.0%;	Pred. No. 1.ee-99;		
	Matches	204; Conservative	0; Mismatches	0; Indels	0; Gaps
OY	394 ccaacatcctggttgccttgagcatgcatactttaactlgaacattgaagtlttcagttcca	453			
Db	414 CCACACCCTCGGTGTCCTGGAGCATGCTCTACTTTTACATGGAAGTTCCAGGTTCCA	355			
OY	454 ggaagccctcagaagcctggcgctgctgtygtcaacctagacaagctgaaggcgctttaaacaag	513			
Db	354 GGAAAGCCCTCAGGCCCTGGGCGTGCTGTGCACCTTACCACCTTAGGAGACTCTTAATACAG	295			
OY	514 aattgctctttagcgacctggaataataataacttaattcgttaaatgatgtaanaacatcta	573			
Db	294 AATTAGCTTTTGTCCACTGGAGCAATAAATTAATTAAATTTGTAAACATGTGAAGAACAATCTA	235			
OY	574 taacatctactgaagcctgcttct	597			
Db	234 TAAACATCTACTGAAGCCTCTTCT	211			

RESULT	5
A1870547/c	
LOCUS	398 bp mRNA EST 07-MAR-2000
DEFINITION	w147a04.x1 NCI_CGAP_Ut1 Homo sapiens CDNA IMAGE:2428013'
	mRNA sequence.
ACCESSION	A1870547
VERSION	A1870547.1 GI:5544515

KEYWORDS	EST.
SOURCE	human.
ORGANISM	Homo sapiens
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS	1 (bases 1 to 398)
TITLE	NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap , National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL	Unpublished (1997)
COMMENT	Contact: Robert Strausberg, Ph.D. Email: craps-r@mail.nih.gov Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
	CDNA Library Preparation: Life Technologies, Inc. CDNA Library Arrayed by: Greg Lennon, Ph.D. DNA Sequencing by: Washington University Genome Sequencing Center Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www.bio.llnl.gov/dbtr/image/image.html Insert length: 1826 Std Error: 0.00 Seq primer: -40UP from Gibco High quality sequence stop: 373. Location/Qualifiers 1. 398 human sapiens
FEATURES	
SOURCE	

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2428014"
/clone_id="NCI-CGAP_0c1"
/tissue_type="well-differentiated endometrial
adenocarcinoma, 7 pooled tumors"
/lab_host="DH10B"
/notes="Organ: uterus; Vector: pCMV-SPORT6; Site_1: Salti.
Site_2: NotI; Cloned unidirectionally. Primer: Oligo dr.
Average insert size 1.75 Kb. life Technologies catalog #:
11538-014"

```

	Dd	398	CTTTGGCAGCTGAGATGAAATTACTTAAATTGTGAACATGCGAATAACATCTATAAACAT	339
	Oy	581	ctaccgaagacctgttcc	597
	Dd	338	CTACTGACCCGTGTCT	322
RESULT	6			
AM979007		269 bp	mRNA	EST
LOCUS				02-JUN-2000
DEFINITION	AM979007			
ACCESION	EST381117	MAGE resequences, MAGP Homo sapiens CDNA, mRNA sequence		
VERSION	AM979007			
KEYWORDS	AM979007.1	GI:8170290		
SOURCE		EST.		
ORGANISM		human,		
		Homo sapiens		
		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
		Eukaryota; Eutheria; Primates; Catarrhini; Hominoidea; Homo.		
		Mammalia; Hominidae; Homo.		
		(bases 1 to 269)		
REFERENCE		Hegde, P., Qi, R., Aternathy, K., Dharap, S., Gaspard, R., Gay, C., Ho,		
AUTHORS		Lee, N.H., Yeatman, T.J., and		

TITLE Quackenhush, J.
Assessment of gene expression patterns in a model of colon tumor metastasis using a 19,200 element cDNA microarray
JOURNAL COMMENT Unpublished (2000)
Contact: John Quackenhush
The Institute for Genomic Research

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9712 Medical Center Dr., Rockville, MD 20850, USA

Tel: 301 838 3528
Fax: 301 838 0208

Email: johngetlgr.org

Plate: 401

Seq primer: Forward.
Location/Qualifiers

1. .269
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone.lib="MAGE_resequencences, MAGE"
/note="Vector: pBluescriptskm"
82 a 69 c 50 g 55 t 13 others

BASE COUNT
ORIGIN

Query Match 7.9%; Score 48; DB 123; Length 269;
Best Local Similarity 100.0%; Pred. No. 7.6e-15; Indels 0; Gaps 0;
Matches 48; Conservative 0; Mismatches 0;

OY 272 ataaataaagacacgtccatccctcccaacgacgcagcagg 319
|||||
DB 113 ATTAATAAAGACACTGCGACGTCCTCCCTCAACGTCGCCAGCCAGG 160

RESULT 7
AT366910/c 234 bp mRNA EST 08-JAN-1999
LOCUS g949409.x1 NCI_CGAP_Ut2 Homo sapiens cDNA clone IMAGE:1989280 3',
DEFINITION mRNA sequence.
ACCESSION AT366910
VERSION AT366910.1 GI:4136655
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens; Chordata; Vertebrata; Euteleostomi;
Eukaryota; Metazoa; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 234)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
Tissue Procurement: Christopher Moskalkuk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: Life Technologies, Inc.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/dbtrp/image/image.html
Seq primer: -400P from Gibco
High quality sequence stop: 226.

FEATURES
source

Location/Qualifiers
1. .234
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1989280"
/clone.lib="NCI_CGAP_Ut2"
/tissue.type="moderately differentiated endometrial
adenocarcinoma, 3 pooled tumors"
/lab_host="DH10B"
/note="Organ: uterus; Vector: PCMV-SPORT6; Site_1: SalI;
Site_2: NotI; Cloned unidirectionally. Primer: Oligo dT.
Average insert size 1.85 kb. Life Technologies catalog #:
11539-012"

BASE COUNT
ORIGIN

54 a 42 c 38 g 100 t

Query Match 4.8%; Score 29; DB 19; Length 234;
Best Local Similarity 100.0%; Pred. No. 0.00016;

Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 550 attgtacaatgtgaacacataaac 578
|||||
DB 122 ATTGTACATGTGAACACATCTATTAAC 94

RESULT 8
AT167719/c 428 bp mRNA EST 20-DEC-1999
LOCUS w339a02.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:2383082 3',
DEFINITION mRNA sequence.
ACCESSION AT167719
VERSION AT167719.1 GI:5234228
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens; Chordata; Vertebrata; Euteleostomi;
Eukaryota; Metazoa; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 428)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
Tissue Procurement: Christopher Moskalkuk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/dbtrp/image/image.html
Insert Length: 276 Std Error: 0.00
Seq primer: -400P from Gibco.

FEATURES
source

Location/Qualifiers
1. .428
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2383082"
/clone.lib="NCI_CGAP_Kid11"
/lab_host="DH10B"
/note="Organ: kidney; Vector: p7T3D-pac (pharmacia) with
a modified polylinker; Site_1: Not I; Site_2: Eco RI;
Plasmid DNA from the normalized library NCI_CGAP_Kid3 was
prepared, and ss circles were made in vitro. Following RAP
purification, this DNA was used as tracer in a subtractive
hybridization reaction. The driver was PCR-amplified cDNAs
from a pool of 5,000 clones made from the same library
(cloneids 1322376-1323911, 1456007-1456775, and
1500552-1502853). Subtraction by Bento Soares and M.
Fatima Bonaldo."

BASE COUNT
ORIGIN

101 a 104 c 57 g 166 t

Query Match 4.1%; Score 25; DB 24; Length 428;
Best Local Similarity 100.0%; Pred. No. 0.024;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 75 gtgggacatgtcactgcatccta 99
|||||
DB 101 GTGGGACATGCTACTGCGACATCTA 77

RESULT 9
AT609595/c 530 bp mRNA EST 14-MAY-1999
LOCUS tw28d08.x1 NCI_CGAP_Ov35 Homo sapiens cDNA clone IMAGE:2261007 3',
DEFINITION mRNA sequence.
ACCESSION AT609595
VERSION AT609595.1 GI:4618762

KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 530)
AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabbs-r@mail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LNLW at:
www-bio.lnl.gov/bbrp/image/image.html
Insert Length: 576 Std Error: 0.00
Seq primer: -400P from Gtbc
High quality sequence stop: 404
POLY-A-No.

FEATURES

1. .530
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2261007"
/clone_lib="NCI-CGAP-OV35"
/tissue_type="tumor, 5 pooled (see description)"
/lab_host="DH10B"
/note="Organ: ovary; Vector: PCMV-SPORT6; Site_1: SalI;
Site_2: NotI; This library represents the normally
version of NCI-CGAP-OV23. Cloned unidirectionally.
Primer: Oligo dT. Average insert size 0.86 kb. Tumor
types include: mixed Mullerian tumor, papillary serous,
clear cell, spindle cell. All are primary tumors,
metastasis positive. Constructed by Life Technologies."

BASE COUNT 131 a 137 c 63 g 198 t 1 others
ORIGIN

Query Match 4.1%; Score 25; DB 22; Length 530;
Best Local Similarity 100.0%; Pred. No. 0.024;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 75 gtgggaacatgctactgcatccta 99
|||||
DB 105 GTGGGAACATGCTACTGCACTCA 81

RESULT 10
LOCUS AW771618 541 bp mRNA EST 04-MAY-2000
DEFINITION hn59c06.x1 NCI-CGAP_Kid11 Homo sapiens cDNA clone IMAGE:3032170 3',
mRNA sequence.
ACCESSION AW771618
VERSION
KEYWORDS
SOURCE EST.
ORGANISM human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 541)
AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabbs-r@mail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.

FEATURES
source
1. .541
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3032170"
/clone_lib="NCI-CGAP_Kid11"
/lab_host="DH10B"
/note="Organ: kidney; Vector: pT73D-Pac (Pharmacia) with
a modified polylinker; Site_1: Not I; Site_2: Eco RI;
Plasmid DNA from the normalized library NCI-CGAP_Kid3 was
prepared, and ss circles were made in vitro. Following HAP
purification, this DNA was used as tracer in a subtractive
hybridization reaction. The driver was PCR-amplified cDNAs
from a pool of 5,000 clones made from the same library
(cloneids 1322376-1323911, 1456007-1456775, and
1500552-1502855). Subtraction by Bento Soares and M.
Fatima Bonaldo."
High quality sequence stop: 463.
Seq primer: -400P from Gtbc
Insert Length: 576 Std Error: 0.00
High quality sequence stop: 463.

BASE COUNT 136 a 136 c 65 g 204 t
ORIGIN

Query Match 4.1%; Score 25; DB 120; Length 541;
Best Local Similarity 100.0%; Pred. No. 0.024;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 75 gtgggaacatgctactgcatccta 99
|||||
DB 101 GTGGGAACATGCTACTGCACTCA 77

RESULT 11
LOCUS BE144594 323 bp mRNA EST 21-JUN-2000
DEFINITION MR0-HT0167-141199-002-C02 HT0167 Homo sapiens cDNA, mRNA sequence.
ACCESSION BE144594
VERSION BE144594.1 GI:8607227
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 323)
AUTHORS Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Brines, M.R.,
Nagel, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,
Goldman, G.H., Carvalhal, A.F., Matsukuma, A., Bala, G.S., Simpson, D.H.,
Brunstein, A., deOliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare,
M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and
Simpson, A.J.
Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
20202663
JOURNAL
MEDLINE
COMMENT Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpone@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(<http://www.ludwig.org.br/scripts/gethtml2.pl?l=612-MR0-HT0167-141>)
199-002-c026t3-1999-11-14&t4=1
Seq primer: puc 18 forward

High quality sequence start: 13
High quality sequence stop: 323.
Location/Qualifiers

FEATURES

SOURCE

```

1. 323
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_id="HT0167"
/dev_stage="Adult"
/Note="Organ: head, neck; Vector: puc18; Site_1: SmaI;
Site_2: SmaI; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters Patent Research)
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the puc 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."
low stringency conditions."
103 a 75 c 79 g 65 t 1 others

```

BASE COUNT

103 a 75 c 79 g 65 t 1 others

ORIGIN

Query Match 3.8%; Score 23; DB 164; Length 323;
Best Local Similarity 100.0%; Pred. No. 0.29;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 163 agaattactagccccaatgctc 185
|||||

Db 135 AGAATTATCTAGCCCAATGCTC 157

RESULT 12

B03713 390 bp DNA GSS 13-JUL-1996
LOCUS B03713
DEFINITION CSRL-185e7-u CSRL flow sorted Chromosome 11 specific cosmid Homo
sapiens genomic clone CSRL-185e7, DNA sequence.

ACCESSION B03713
B03713.1 GI:1412991

KEYWORDS

SOURCE

ORGANISM

Human.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 390)
Evans, G.A., Burbee, D., Davies, C., Hahner, L., Oliver, T., Gilbert, M.,
Jones, D., Ward, T., Gillilan, E., Schagemann, J., Probst, S., Harris
, J., Depford, J., McFarland, J., Burzinski, K., Khan, M., Kupfer, K. and
Garner, H.R.

Genomic Sequence Sampled Map of Chromosome 11
Unpublished (1996)

TITLE

JOURNAL

COMMENT

Contact: Evans GA, Shane Probst
McMort Center for Human Growth and Development
University of Texas Southwestern Medical Center At Dallas
5323 Harry Hines Blvd, Dallas TX 75235-8591
Tel: 214-648-1600
Fax: 214-648-1666
Email: g.evans@uts.wmed.edu, shane@mcdermott.wmed.edu
PCR Primers
FORWARD: ACAAGAAATATCTAGCCCC
BACKWARD: AACCAACAACCTCACACAC
Seq primer: 77
Class: cosmid ends
High quality sequence stop: 390.
Location/Qualifiers

FEATURES

SOURCE

```

1. 390
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_id="CSRL-185e7"
/clone_lib="CSRL flow sorted Chromosome 11 specific
cosmid"
/sex="female"
/cell_type="chimeric hamster somatic cell hybrid"
/Note="Vector: scos-1; Human Chromosome 11 specific cosmid
library prepared from flow sorted human Chromosome 11
derived from Chinese Hamster Ovary (CHO) monochromosomal
somatic cell hybrid, J1"

```

BASE COUNT 85 a 111 c 70 g 113 t 11 others
ORIGIN

Query Match 3.8%; Score 23; DB 256; Length 390;
Best Local Similarity 100.0%; Pred. No. 0.29;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 163 agaattactagccccaatgctc 185
|||||

Db 51 AGAATTATCTAGCCCAATGCTC 73

RESULT 13

AQ282338/c 528 bp DNA GSS 27-APR-1999
LOCUS AQ282338
DEFINITION RPI11-79011.TV RPI1-11 Homo sapiens genomic clone RPI1-11-79011,
DNA sequence.

ACCESSION AQ282338
AQ282338.1 GI:3908887

KEYWORDS

SOURCE

Human.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 528)
Adams, M.D., Rounsley, S.D., Zhao, S., Bass, S., Linher, K., Golden, K.,
Berry, K., Granger, D., Suh, E., Wible, C., de Jong, P. and Ventier, J.C.

Use of human BAC End Sequences for Sequence-Ready Map Building
Unpublished (1998)

TITLE

JOURNAL

COMMENT

Contact: Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: mdamas@tigr.org
Clones are derived from the human BAC library RPI1-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased or from
BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from
Research Genetics (info@resgen.com). BAC end search page:
http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html
Seq primer: 47
Class: BAC ends.
Location/Qualifiers

FEATURES

SOURCE

```

1. 528
/organism="Homo sapiens"
/db_xref="GDB:7530298"
/db_xref="taxon:9606"
/clone="RPI1-11-79011"
/clone_lib="RPI1-11"
/sex="Male"
/cell_type="Lymphocytes"
/Note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
RPI11 Human Male BAC library"

```

BASE COUNT

162 a 96 c 135 g 135 t

ORIGIN

Query Match 3.8%; Score 23; DB 226; Length 528;
Best Local Similarity 100.0%; Pred. No. 0.29;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 163 agaattactagccccaatgctc 185
|||||

Db 500 AGAATTATCTAGCCCAATGCTC 478

RESULT 14

R77087 422 bp mRNA EST 06-JUN-1995
LOCUS R77087
DEFINITION y164904.r1 Soares placenta ND2HP Homo sapiens cDNA clone

Wed Nov 7 09:21:19 2001

us-09-656-668-198.oli.rst

IMAGE:144054 5' similar to contains LTR4 repetitive element ;, mRNA sequence.

ACCESSION R77087
 VERSION R77087.1 GI:851719
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 422)
 Hillier L., Clark N., Dubuque T., Elliston K., Hawkins M., Holman M., Hultman M., Kucaba T., Le M., Lennon G., Marra M., Parsons J., Rikkin L., Rohlfing T., Soares M., Tan F., Trevisakis E., Waterston R., Williamson R., Wohldmann P. and Wilson R.
 The Mashu-Merck EST Project
 Unpublished (1995)
 Contact: Milson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 Insert Size: 587
 High quality sequence stops: 333
 Source: IMAGE Consortium, LNL
 This clone is available royalty-free through LNL ; contact the IMAGE Consortium (info@image.lnl.gov) for further information.
 Putative full length read
 Insert length: 587 Std Error: 0.00
 Seq primer: M13Kp1
 High quality sequence stop: 333.
 Location/Qualifiers

FEATURES
 source
 1..422
 /organism="Homo sapiens"
 /db_xref="GDB:553380"
 /db_xref="taxon:9606"
 /clone="IMAGE:144054"
 /clone_id="Soares placenta ND2HP"
 /sex="Female"
 /dev_stage="placenta obtained at birth (full term)"
 /lab_host="DH10B (ampicillin resistant)"
 /note="Organ: placenta; Vector: pT7p3d (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5', AACTGGAAGATTCGGCGGCGAGATTTTCTTTT 3'], (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7p3 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 102 a 93 c 88 g 134 t 5 others

ORIGIN

Query Match 3.5%; Score 21; DB 188; Length 422;
 Best Local Similarity 100.0%; Pred. No. 3.5;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 245 tgccctgtgtcgtgctctgt 265
 ||||||||||||||||||
 Db 350 tgccttctgtcgtcgtctct 370

RESULT 15
 AL513616 1018 bp mRNA EST 13-FEB-2001
 LOCUS AL513616 LTI_NFL006_PL2 Homo sapiens cDNA clone XCL08B0012E07 5
 DEFINITION prime, mRNA sequence.
 ACCESSION AL513616
 VERSION AL513616.1 GI:12777110
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 1018)
 Li W.B., Gruber C., Jesssee J. and Polayes D.
 Full-length cDNA libraries and normalization
 Unpublished (2001)
 TITLE JOURNAL
 COMMENT Contact: Genoscope
 Genoscope - Centre National de Sequencage
 BP 191 91006 EVRY cedex - France
 Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr.
 Location/Qualifiers

FEATURES
 source
 1..1018
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="XCL08B0012E07"
 /clone_id="LTI_NFL006_PL2"
 /issue_type="placenta"
 /note="Vector: pCMVSPORT 6; Site_1: NotI; 1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end was enriched, double-stranded cDNA was digested with Not I and cloned into the Not I and Eco RV sites of the pCMVSPORT 6 vector. Library was normalized. Library was constructed by Life Technologies. Contact : Peng Liang Life Technologies, a division of Invitrogen 9800 Medical Center Drive Rockville, Maryland 20850, USA Fax : (1) 301 610 8371
 Email : filiang@lifetech.com URL : http://fulllength.invitrogen.com" 4 others

BASE COUNT 212 a 308 c 318 g 176 t 4 others

ORIGIN

Query Match 3.5%; Score 21; DB 105; Length 1018;
 Best Local Similarity 100.0%; Pred. No. 3.7;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 459 gccctcagcctgagctgctg 479
 ||||||||||||||||||
 Db 31 gccctcagcctcagcctcgt 11

Search completed: November 5, 2001, 23:19:10
 Job time: 2595 sec

Wed Nov 7 09:21:19 2001

us-09-656-668-198.oli.rst

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

W nucleic - nucleic search, using sw model

run on: November 5, 2001, 22:16:10 ; Search time 1182.83 Seconds
(without alignments)
7924.601 Million cell updates/sec

title: us-09-656-668-198

effective score: 606
1 tgagttgcccccttaccac.....aagcctgttctgtctgcac 606

sequence: IDENTITY_NUC
Gapop 10.0 / Gapext 1.0

coring table: 1344157 seqs, 7733874588 residues

searched: 2688314

total number of hits satisfying chosen parameters:

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : GenBank:
1: gb_ba1:*
2: gb_ba2:*
3: gb_ba3:*
4: gb_in1:*
5: gb_in2:*
6: gb_in3:*
7: gb_cm:*
8: gb_ov:*
9: gb_pat1:*
10: gb_pat2:*
11: gb_ph:*
12: gb_pl1:*
13: gb_pl2:*
14: gb_pl3:*
15: gb_pl4:*
16: gb_ba1:*
17: gb_ba2:*
18: gb_fun:*
19: em_htgo_hum:*
20: em_htgo_inv:*
21: em_htgo_rod:*
22: em_htgo_hum1:*
23: em_htgo_hum2:*
24: em_htgo_hum3:*
25: em_htgo_hum4:*
26: em_htgo_hum5:*
27: em_htgo_hum6:*
28: em_htgo_hum7:*
29: em_htgo_hum8:*
30: em_htg_hum1:*
31: em_htg_hum2:*
32: em_htg_rod:*
33: em_htg_rod:*
34: em_hum1:*
35: em_hum2:*
36: em_hum3:*
37: em_hum4:*
38: em_hum5:*
39: em_hum6:*
40: em_hum7:*
41: em_in:*
42: em_cm:*
43: em_or:*

44: em_ov:*
45: em_pat:*
46: em_ph:*
47: em_pl:*
48: em_ro:*
49: em_sts:*
50: em_sy:*
51: em_un:*
52: em_v1:*
53: gb_sts1:*
54: gb_sts2:*
55: gb_sts3:*
56: gb_sy:*
57: gb_un:*
58: gb_v1:*
59: gb_v12:*
60: gb_v12:*
61: gb_htg1:*
62: gb_htg2:*
63: gb_htg3:*
64: gb_htg4:*
65: gb_htg5:*
66: gb_htg6:*
67: gb_htg7:*
68: gb_htg8:*
69: gb_htg9:*
70: gb_htg10:*
71: gb_htg11:*
72: gb_htg12:*
73: gb_htg13:*
74: gb_htg14:*
75: gb_htg15:*
76: gb_htg16:*
77: gb_htg17:*
78: gb_htg18:*
79: gb_htg19:*
80: gb_htg20:*
81: gb_htg21:*
82: gb_htg22:*
83: gb_htg23:*
84: gb_htg24:*
85: gb_htg25:*
86: gb_pr1:*
87: gb_pr2:*
88: gb_pr3:*
89: gb_pr4:*
90: gb_pr5:*
91: gb_pr6:*
92: gb_pr7:*
93: gb_pr8:*
94: gb_pr9:*
95: gb_ro1:*
96: gb_ro2:*
97: gb_in4:*
98: gb_pr10:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	606	100.0	606	10	AX093380 Sequence
2	104.6	17.3	144014	66	AC021763 Homo sapi
3	104.6	17.3	159150	83	AP002438 Homo sapi
4	104.6	17.3	187200	82	AP001379 Homo sapi
5	97.2	16.0	152081	86	AC007381 Homo sapi
6	95.6	15.8	93714	79	AL161661 Homo sapi
7	95.6	15.8	147815	92	AL031687 Human DNA
8	93.6	15.4	185143	69	AC025861 Homo sapi

9	93.6	15.4	208763	79	AL153791	AL153791 Homo sapi
10	91.8	15.1	147929	87	AC016951	AC016951 Homo sapi
9	91.8	15.0	194313	79	AL153759	AL153759 Homo sapi
11	91	15.0	207746	64	AC011708	AC011708 Homo sapi
12	91	15.0	162082	66	AC021051	AC021051 Homo sapi
13	90.8	15.0	180861	74	AC073244	AC073244 Homo sapi
14	90.8	15.0	180861	66	AC073244	AC073244 Homo sapi
15	90.4	14.9	63020	73	AC068945	AC068945 Homo sapi
16	90.4	14.9	195141	73	AC022612	AC022612 Homo sapi
17	89.6	14.8	163511	90	AL442203	AL442203 Human DNA
18	89.6	14.8	177962	71	AC027806	AC027806 Homo sapi
19	89.6	14.8	186133	79	AL157932	AL157932 Homo sapi
20	89.6	14.8	191831	66	AC021149	AC021149 Homo sapi
21	89.6	14.8	325013	62	AC016817	AC016817 Homo sapi
22	89.2	14.7	155375	94	HS914914	AL103117 Human DNA
23	88.6	14.6	160841	81	AL449323	AL449323 Homo sapi
24	88.4	14.6	160421	69	AC025005	AC025005 Homo sapi
25	88.4	14.6	160840	79	AL153766	AL153766 Homo sapi
26	88.2	14.6	142796	86	AC005844	AC005844 Homo sapi
27	88.2	14.6	142796	86	AC005844	AC005844 Homo sapi
28	88.2	14.6	182810	67	AL354914	AL354914 Homo sapi
29	87.8	14.5	189072	71	AC022491	AC022491 Homo sapi
30	87.6	14.5	168402	73	AC0040965	AC0040965 Homo sapi
31	87	14.4	107462	73	AC062715	AC062715 Homo sapi
32	87	14.4	189351	73	AC068588	AC068588 Homo sapi
33	87	14.4	193577	80	AC068025	AC068025 Homo sapi
34	86.8	14.3	110000	73	AL1390202_07	Continuation (8 of
35	86.8	14.3	172096	65	AC079864	AC079864 Homo sapi
36	86.8	14.3	218708	66	AC021166	AC021166 Homo sapi
37	86.2	14.2	133702	90	AL139182	AL139182 Human DNA
38	86.2	14.2	153178	61	AC009929	AC009929 Homo sapi
39	86.2	14.2	168293	65	AC0018397	AC0018397 Homo sapi
40	85.8	14.2	115564	92	HS1170D6	AL103095 Human DNA
41	85.8	14.2	181218	71	AC036222	AC036222 Homo sapi
42	85.8	14.2	184834	88	AC026122	AC026122 Homo sapi
43	85.6	14.1	146652	81	AL445504	AL445504 Homo sapi
44	85.6	14.1	151357	62	AC011573	AC011573 Homo sapi
45	85.6	14.1	157912	89	AL133387	AL133387 Human DNA

ALIGNMENTS

[illegible]

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	Conservative	0;			
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Db	121	TGCTAAACCTCTTTCAACGCACAGCAGCAAGCCCCAAMAAAGAAATTATCTACCCCA	180
OY	181	atgtccataaacactgctgtgtgagaaacctaacccggaagctcttactggtcttctaa	240
Db	181	ATGTCCATTAACCTGCTGTGAGAAACCTACCCGAGATCTTACTGGGCTTCATAGSTA	240
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Db	241	AGCTTGCCCTTGTCTGTGGCTTCGTGAATATATATAAATAAGACCTGCCAGTCCCTCC	300
OY	301	ctcacaagctcccgagccagggcttcaaggaacatttccaataacagtagaatgaacactaata	360
Db	301	CTCAACGCTCCGAGCCAGGAGGCTCAAGGCAATTCGAATAGAGTGAATGACACTTAATA	360
OY	361	tgtatttcaaatctcagcaactagaaatgacaaacacatccgtgttgctctggagctg	420
Db	361	TGTATTTCAAAATCTCAGCACTAGAGATGACCAACATCTGTTGGCTCGGAGACTG	420
OY	421	tccatgattttagcaattgaaagtcttaggttccaggaagccctcaaggctggctgcgcgg	480
Db	421	TCCATGATTTCAGCAATTGAAAGTTTCAGGTTTCAGAGAAACCTCAGCGCTGGCTCTGG	480
OY	481	tcaacctcagcagctgagagacatcttcaatacagaatagtcttcttgactggagatga	540
Db	481	TCACCTTACAGCTGAGGAGCTTCTTCATATACAGAAATTAGCTTTTGACATGGAGATGA	540
OY	541	tataacttaatttgtaacatgtagaaacatctataaacaatcactcgaagcctgttctgt	600
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OY	601	ctgcac 606	
Db	601	CTGCAC 606	

RESULT	2	13-OCT-2000
ACLOCUS	AC021763/C	
LOCUS	AC021763	144014 bp
DEFINITION	Homo sapiens chromosome 18 clone Rpl1-56021 map 18,	WORKING DRAFT
ACCESSION	AC021763	
VERSION	AC021763.3	GI:10800280
KEYWORDS	HTG: HTGS_PHASE1; HTGS_DRAFT.	
SOURCE	human.	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.	
AUTHORS	1 (bases 1 to 144014)	
JOURNAL	2 (bases 1 to 144014)	
REFERENCE	Unpublished	
AUTHORS	Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N., Anderson,S., Baldwin,J., Barna,N., Beckelly,R., Beda,F., Boguslavsky,L., Boukhalter,B., Brown,A., Burnett,G., Castle,A., Choepel,Y., Colangelo,M., Collins,S., Collamore,A., Cooke,P., Dearellano,K., Dewar,K., Domino,M., Doyle,M., Fenesstor,J., Ferreira,P., Flitzhugh,W., Forrest,C., Gage,D., Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heatord,A., Karatas,A., Klein,J., Howland,J.C., Lehoczy,J., Levine,C., Jones,C., Kann,L., Karas,A., McKernan,K., McDonald,P., Merquis,N., Meneus,L., Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P., Oliveira,T.M., Peterson,K., Pierre,N., Pisanic,C., Pollara,V., Raymond,C., Riley,R., Rothman,D., Roy,A., Santos,R., Severy,P., Spencer,B., Stange-Thomann,N.,	

us-09-656-668-198.rge

Wed Nov 7 09:21:19 2001

VERSTON
 KEYWORDS
 SOURCE
 ORGANISM

AP002438.1 GI:8307742
 HTG: HTGS-PHASE1; HTGS-DRAFT.
 Homo sapiens DNA, clone: RP11-679N11.
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
 AUTHORS

1 (bases 1 to 159150)
 Hattoni, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
 Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 Homo sapiens 159,150 genomic DNA of 18q21
 Published Only in DataBase (2000) In press

TITLE
 JOURNAL

REFERENCE
 AUTHORS

2 (bases 1 to 159150)
 Hattoni, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
 Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 Direct Submission
 Submitted (02-JUN-2000) Masahira Hattoni, The Institute of Physical
 and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
 Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
 Japan (E-mail: hattoni@gsc.riken.go.jp,
 URL: http://hgp.gsc.riken.go.jp/, Tel: 81-42-778-9923,
 Fax: 81-42-778-9924)

COMMENT

----- Genome Center
 Center: RIKEN Genomic Sciences Center (GSC)
 Center code: RIKEN
 Web site: http://hgp.gsc.riken.go.jp/
 Contact: hattoni@gsc.riken.go.jp
 ----- Project Information
 Center project name: Humdraft18
 Center clone name: RP11-679N11
 ----- Summary Statistics
 Sequencing vector: PCR products; 100% of reads
 Chemistry: Dye-terminator ET-amersham; 100% of reads
 Assembly program: Phrap; version 0.990329
 Consensus quality: 141170 bases at least Q40
 Consensus quality: 150107 bases at least Q30
 Consensus quality: 153808 bases at least Q20
 Insert size: 155950; sum-of-contigs
 Quality coverage: 4.52x in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of 33 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1
 18093 contig of 18093 bp in length
 18194 34028 contig of 15835 bp in length
 34129 47853 contig of 13725 bp in length
 47954 61375 contig of 13422 bp in length
 61476 77424 contig of 100 bp
 69864 77424 contig of 100 bp
 77525 83516 contig of 5992 bp in length
 83617 89288 contig of 5672 bp in length
 89389 94381 contig of 4993 bp in length
 94382 94481 contig of 100 bp
 94482 100220 contig of 5739 bp in length
 100221 100320 contig of 4737 bp in length
 100321 105057 contig of 100 bp
 105058 105157 contig of 100 bp
 105158 110565 contig of 5408 bp in length
 110566 110665 contig of 100 bp
 110666 115962 contig of 5297 bp in length
 115963 116062 contig of 100 bp
 116063 119619 contig of 3557 bp in length
 119620 119719 contig of 100 bp
 119720 124300 contig of 4581 bp in length
 124301 124400 contig of 100 bp
 124401 127169 contig of 2769 bp in length
 127170 127269 contig of 100 bp
 127270 129941 contig of 2672 bp in length
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 130042 132701 contig of 2660 bp in length
 132702 132801 contig of 100 bp
 132802 135223 contig of 2422 bp in length
 135224 135323 contig of 100 bp
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 139041 140352 contig of 1312 bp in length
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 140453 142867 contig of 2415 bp in length
 142868 142967 contig of 100 bp
 142968 144444 contig of 1477 bp in length
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 144545 146121 contig of 1577 bp in length
 146122 146221 gap of 100 bp
 146222 147660 contig of 1439 bp in length
 147661 147760 gap of 100 bp
 147761 149057 contig of 1297 bp in length
 149058 149157 gap of 100 bp
 149158 150648 contig of 1491 bp in length
 150649 150748 gap of 100 bp
 150749 152399 contig of 1651 bp in length
 152400 152499 gap of 100 bp
 152500 154103 contig of 1604 bp in length
 154104 154203 gap of 100 bp
 154204 155806 contig of 1603 bp in length

AP001379 187200 bp DNA HTG 15-JUL-2000
 LOCUS Homo sapiens chromosome 18 clone RP11-850H3 map 18q12, WORKING
 DEFINITION DRAFT SEQUENCE, 21 unordered pieces.
 AP001379 3 GI-9229958
 ACCESSION HTG: HTGS_PHASE1; HTGS_DRAFT.
 VERSION DRAFT SEQUENCE, 21 unordered pieces.
 KEYWORDS Homo sapiens DNA, clone:RP11-850H3.
 SOURCE Homo sapiens
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 187200)
 AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P.,
 Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 TITLE Homo sapiens 187,200 genomic DNA of 18q12
 JOURNAL Published Only in DataBase (2000) In press
 REFERENCE 2 (bases 1 to 187200)
 AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T. D., Hong-Seog, P.,
 Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 TITLE Direct Submission
 JOURNAL Submitted (09-MAR-2000) Masahira Hattori, The Institute of Physical
 and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555, Japan
 (E-mail:hattori@gsc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/,
 Tel:81-42-778-9923, Fax:81-42-778-9924)
 COMMENT On Jul 15, 2000 this sequence version replaced gi:8117792.
 ----- Genome Center
 Center: RIKEN Genomic Sciences Center(GSC)
 Center code: RIKEN
 Web site: http://hgp.gsc.riken.go.jp/
 Contact: hattori@gs.riken.go.jp
 ----- Project Information
 Center project name: HumDraft18
 Center clone name: RP11-850H3
 ----- Summary Statistics
 Sequencing vector: PCR products; 100% of reads
 Chemistry: Dye-terminator ET-amersham; 100% of reads
 Assembly program: Phrap; version 0.990329
 Consensus quality: 179971 bases at least Q40
 Consensus quality: 182832 bases at least Q30
 Consensus quality: 184194 bases at least Q20
 Insert size: 185200; sum-of-contigs
 Quality coverage: 9.46x in Q20 bases; sum-of-contigs

 NOTE: This is a 'working draft' sequence. It currently consists of
 21 contigs. The true order of the pieces is not known and their
 order in this sequence record is arbitrary. Gaps between the
 contigs are represented as runs N, but the exact sizes of the gaps
 are unknown. This record will be updated with the finished sequence
 as soon as it is available and the accession number will be
 preserved
 1 33581 contig of 33581 bp in length
 33682 58093 contig of 24412 bp in length
 58194 77625 contig of 19432 bp in length
 77726 97031 contig of 19306 bp in length
 97132 111990 contig of 14859 bp in length
 112091 125465 contig of 13375 bp in length
 125566 144628 contig of 9889 bp in length
 135555 151062 contig of 9074 bp in length
 14729 155111 contig of 6334 bp in length
 151163 160186 contig of 4349 bp in length
 155612 164088 contig of 4575 bp in length
 160287 167739 contig of 3802 bp in length
 164189 171188 contig of 3551 bp in length
 167840 174981 contig of 3349 bp in length
 171289 177828 contig of 3693 bp in length
 175082 177929 contig of 2747 bp in length
 177929 180660 contig of 2732 bp in length
 180761 183192 contig of 2432 bp in length
 183293 185028 contig of 1736 bp in length
 185129 185875 contig of 747 bp in length
 185976 187200 contig of 1225 bp in length
 * NOTE: This is a 'working draft' sequence. It currently

* NOTE: This is a 'working draft' sequence. It currently

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* 155907 155906: gap of 100 bp
* 155907 157204: contig of 1298 bp in length
* 157205 157304: gap of 100 bp
* 157305 158735: contig of 1431 bp in length
* 158736 158835: gap of 100 bp
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17.38; Score 104.6; DB 83; Length 159150;
Query Match
Best Local Similarity 64.94; Pred. No. 6.7e-20;
Matches 155; Conservative 0; Mismatches 84; Indels 0; Gaps 0;

QY 24 ccagtgaaatttgcaattcctaataagacgtgttttgattcacacctgggtgggaac 83
||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
DB 24951 CCCAGTAATAATTGGCAATGCCCGAGGACATTTTGGTTGTCACACGGGAACAGGAC 25010
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QY 84 atgtactggcatctaagtcatagggcgaagtgtctgctgaacatctttcaacgcaca 143
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DB 25011 CTGCTACTGGCACTTAATGAGTAGGATGGCTGCTGAACATCTCTACAGTGCACA 25070
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QY 144 ggacagagcccccacaaagagaattatctagcccaaatgtccataacactgctgttag 203
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DB 25071 GGTGAGAACCCCATACAAAGAAGTATGATCCAGCCCAATAATATCACTAGTGGCGAGGCTGAG 25130
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||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
DB 25131 AAACCTGTTCTAAGGCATTTGCAAGTAGCCATCCATGAGAGTGATGCTGTGAAGACGTGTTTC 25189
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* consists of 21 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* * be preserved.
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* 33581: contig of 33581 bp in length
* 33582 33681: gap of 100 bp
* 33682 58093: contig of 24412 bp in length
* 58094 58193: gap of 100 bp
* 58194 77625: contig of 19432 bp in length
* 77626 77725: gap of 100 bp
* 77726 97031: contig of 19306 bp in length
* 97032 97131: gap of 100 bp
* 97132 111990: contig of 14859 bp in length
* 111991 112090: gap of 100 bp
* 112091 125465: contig of 13375 bp in length
* 125466 125565: gap of 100 bp
* 125566 133454: contig of 9889 bp in length
* 133455 133554: gap of 100 bp
* 133555 144628: contig of 9074 bp in length
* 144629 144728: gap of 100 bp
* 144729 151062: contig of 6334 bp in length
* 151063 151162: gap of 100 bp
* 151163 155511: contig of 4349 bp in length
* 155512 155611: gap of 100 bp
* 155612 160186: contig of 4575 bp in length
* 160187 160286: gap of 100 bp
* 160287 164088: contig of 3802 bp in length
* 164089 164188: gap of 100 bp
* 164189 167739: contig of 3551 bp in length
* 167740 167839: gap of 100 bp
* 167840 171188: contig of 3349 bp in length
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* 171289 174981: contig of 3693 bp in length
* 174982 175081: gap of 100 bp
* 175082 177828: contig of 2747 bp in length
* 177829 177928: gap of 100 bp
* 177929 180660: contig of 2732 bp in length
* 180661 180760: gap of 100 bp
* 180761 183192: contig of 2432 bp in length
* 183193 183292: gap of 100 bp
* 183293 185028: contig of 1736 bp in length
* 185029 18528: gap of 100 bp
* 185129 185875: contig of 747 bp in length
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FEATURES
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185976. .187200
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ORIGIN

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BASE COUNT 51230 a 39996 C 41403 9 32571 5
ORIGIN

Query Match	17.3%	Score 104.7	0	Gaps
Best Local Similarity	64.9%	Pred. No. 6.8e-20		
Matches	155; Conservative	0; Mismatches	84; Indels	0; Gaps
QY	24	ccacgtaaatattgcgaattcctaagaagcgtgtttttgattgtcacacctgggtgggaac	83	
		CCCCGATAAATATTTTGGCAATGCCCGAGGACATTTTGGTTGTCACACGGGAACAGGNAC	116087	
Db	116028	CCCCGATAAATATTTTGGCAATGCCCGAGGACATTTTGGTTGTCACACGGGAACAGGNAC	116087	
QY	84	atgctactggtcatcctaagtcatagaggcggaagtaatgctgctaaacatcttttcaacgcaca	143	
		TT	116147	
Db	116088	CTGTACTGTCATCTAATGAGTAGAGGTGAAGGATGCTGCTGACATCTCTACAGTGCACA	116147	
QY	144	ggacagagccccacaaaagaagaattatctagcccaaatgtccataacactgctgttaag	203	
		TT	116207	
Db	116148	GGTCAGAACCCCATACAAAGAAGTATCCAGCCCAAAAATATCACTAGTGCAGAGGCTGAG	116207	
QY	204	aaaacctaccgcaggaatcttactgggttcttaaggtaagcttgcctttgtcttggtcttc	262	
		TT	116266	
Db	116208	AAACCTGTTCTTAGGGCATTTGCCAAAGTACATGAGATGATCTCTGTAAGAGCTGGTTTC	116266	

RESULT	5				PRI	30-SEP-2000
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DEFINITION		Homo sapiens BAC clone RP11-158121	from 2,	complete sequence.		
ACCESSION		AC007381				
VERSION		AC007381.3	GI:6604532			
KEYWORDS		HTG.				
SOURCE		human.				
SCORE						

SOURCE	ORGANISM	REFERENCE	AUTHORS	TITLE	JOURNAL	MEDLINE
human.	Human sapiens					
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.						
1 (bases 1 to 152081)						
Sulston, J.E. and Waterston, R.						
Toward a complete human genome sequence						
Genome Res. 8 (11), 1097-1108 (1998)						
99063792						
2 (bases 1 to 152081)						
Kozlowicz, A., Wohlmann, P. and Harper, M.						
The sequence of Homo sapiens BAC clone RP11-158121						
Unpublished						
3 (bases 1 to 152081)						
Waterston, R.H.						
Direct Submission						
Submitted (25-APR-1999)						
Genome Sequencing Center, Washington						

Wed Nov 7 09:21:19 2001

University School of Medicine, 4444 Forest Park Parkway, St. Louis,

REFERENCE
AUTHORS

MO 63108, USA
4 (bases 1 to 152081)
Waterston, R.H.
Direct Submission
Submitted (19-DEC-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA

REFERENCE
AUTHORS

5 (bases 1 to 152081)
Waterston, R.H.
Direct Submission
Submitted (21-DEC-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA

REFERENCE
AUTHORS

6 (bases 1 to 152081)
Waterston, R.
Direct Submission
Submitted (30-SEP-2000) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Dec 20, 1999 this sequence version replaced gi:5001522.

COMMENT

Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc>
Contact: sapiens@wustl.wustl.edu
----- Summary Statistics

Center project name: H_NH0158121

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Woon, P.Y., Zhao, B., Frengen, E., Tatenno, M., Catanese, J.J. and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBACE3.6
NEIGHBORING SEQUENCE INFORMATION:
The clone sequenced to the right is RP11-119H15. Actual start of this clone is at base position 1 of RP11-158I21.

The sequence RP11-158I21 contains a dinucleotide (CT) repeat from base positions 6270 to 7430 in which the exact length is unknown. A PCR product from clone DNA confirms the approximate size of the region with band sizes of 2527 real, 2508 insilico for EcoRV; and 6293 real, 6234 insilico for HindIII.

FEATURES
SOURCE

Location/Qualifiers
1. 152081
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/db_xref="taxon:9606"
/chromosome="2"
/map="2"

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	/clone_lib="RPCI-11"	
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repeat_region	1252..1426	/note="match to EST N54976 (NID:g1196296) yv38h03.s1"
repeat_region	2507..2687	/rpt_family="(CA)n"
repeat_region	3468..3596	/rpt_family="MER1_type"
repeat_region	3631..3662	/rpt_family="MIR"
repeat_region	5619..5723	/rpt_family="(CAAAA)n"
misc_feature	5619..5723	/note="similar to Mus musculus EST AI255229 (NID:g3862754)
misc_feature	5637..5723	u132h10.y1
misc_feature	5637..5723	/note="similar to Mus musculus EST AA254455 (NID:g1889058)
misc_feature	5641	val6e12.rl
misc_feature	5641	/note="match to EST R88177 (NID:g946990) ym90e05.rl"
misc_feature	5669..5723	/note="similar to Mus musculus EST AW123279 (NID:g6098809)"
misc_feature	5684..5723	/note="match to EST AI081350 (NID:g3418142) ox76e08.x1"
misc_feature	5698..5716	/note="match to EST R88177 (NID:g946990) ym90e05.rl"
repeat_region	6293..7016	/rpt_family="(CCA)n"
repeat_region	7023..7383	/rpt_family="(CCA)n"
repeat_region	8687..8848	/rpt_family="MIR"
repeat_region	8934..8959	/rpt_family="(TAAA)n"
repeat_region	11635..11696	/rpt_family="MIR"
repeat_region	11845..11891	/rpt_family="L2"
misc_feature	12021..12411	/note="match to EST R88177 (NID:g946990) ym90e05.rl"
misc_feature	12031..12173	/note="similar to Mus musculus EST AI255229 (NID:g3862754)
misc_feature	12031..12173	u132h10.y1
misc_feature	12031..12173	/note="match to EST AI081350 (NID:g3418142) ox76e08.x1"
misc_feature	12031..12173	/note="similar to Mus musculus EST AW123279 (NID:g6098809)"
misc_feature	12031..12173	/note="similar to Mus musculus EST AA254455 (NID:g1889058)
misc_feature	12031..12173	val6e12.rl
misc_feature	12073..12173	/note="similar to Mus musculus EST AA183217 (NID:g1766873)
misc_feature	12091..12173	/note="similar to Mus musculus EST AA002793 (NID:g1446526)
misc_feature	12101..12173	mq41q01.rl
misc_feature	12101..12173	/note="similar to Mus musculus EST AA726437 (NID:g2744144)
misc_feature	12117..12173	vu92b11.rl
misc_feature	12124..12173	/note="match to EST AI560419 (NID:g4510760) tn09h07.x1"
misc_feature	12124..12173	/note="similar to Mus musculus EST AA003295 (NID:g1446732)
misc_feature	12134..12173	mq48f11.rl
misc_feature	12598..13767	/note="match to EST AA236279 (NID:g1858428) zr54f05.s1"
misc_feature	12974..13396	/note="CpG island (80C=66.2, o/e=0.80, #CpGs=110)"
repeat_region	12983..13055	/note="match to EST AW157755 (NID:g6229156) au86d06.x1"

Center: Sanger Centre
Center code: SC
Web site: <http://www.sanger.ac.uk>
Contact: humquery@sanger.ac.uk
----- Project Information
Center project name: dj984123

```

----- Summary Statistics -----
Assembly program: XGAP4; version 4.5
Sequencing vector: plasmid; L08752; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Consensus quality: 93681 bases at least Q40
Consensus quality: 93706 bases at least Q30
Consensus quality: 93713 bases at least Q20
Insert size: 93714; sum-of-contigs
Insert size: 109355; 1.0% error; agarose-fp
Quality coverage: 8.25x in Q20 bases; sum-of-contigs
Quality coverage: 8.020x in Q20 bases; agarose-fp

```

* NOTE: This is a 'working draft' sequence.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

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FEATURES
  source
    Location/Qualifiers
      1..93714
        /organism="Homo sapiens"
        /db_xref="taxon:9606"
        /chromosome="20"
        /clone="RP5-984I23"
        /clone_lib="RPCI-5"
      1..93714
        misc_feature
          /note="assembly fragment:00029"
          32669..24035 t

```

BASE COUNT	24148 a	22562 c	22969 g	24035 e
ORIGIN				

Query Match	15.8%;	Score 95.6;	DB 79;	Length 93714;
Best Local Similarity	75.8%;	Pred. No. 2.9e-17;		
Matches 144;	Conservative	0;	Mismatches 44;	Indels 2;
Gaps				
20	ccatcccaagtgaatatgtgcaatctctaaagacgtgttttgattgtcacaacctggtggtg	79		
QY				
44680	CCACCACGGGACATTTACCAATGACTAGAGATGT-TTCTGTGTGCACAACCTGGGT-GG	44623		
DB				
80	gaacatgctactgcatctctaatgcataagagggcagtaatgctgctaaacatcttttcaacg	139		
QY				
44622	GAGGTGCTGCTGGCATCTAGTGGGTGAGGCCAGGAATGCTGCTTAACATCTTACAATG	44563		
DB				
140	cacagcagcagagcccccacaaagagaatattctagccccaatgctccataaacactgctgt	199		
QY				
44562	CCCGGGACAGCCCTCCACGACAAAGAAATTATCCAGGCCCCCAATGTCAATAGAGCTGAGGT	44503		
DB				
200	tgagaaaaacc	209		
QY				
DB	44502	TGAGAAACC	44493	

RESULT	7	PRI	04-APR-2001
H5998H6		DNA	
LOCUS	H5998H6	147815 bp	
DEFINITION	Human DNA sequence from clone RP5-998H6 on chromosome 20q13.1. Contains the gene for the ortholog of rat PB-Cadherin, ESRs, STSS, GGS, two CpG islands and genomic marker D20S17, complete sequence.		
ACCESSION	AL031687		
VERSION	AL031687.17	GI:13366276	
KEYWORDS	htg; CpG island; D20S17; PB-Cadherin.		
SOURCE	human.		
ORGANISM	Homo sapiens		
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
REFERENCE	1 (bases 1 to 147815)		
AUTHORS	Laird,G.		
TITLE	Direct Submission		
JOURNAL	Submitted (27-MAR-2001) Sanger Centre, Hinxton, Cambridgeshire,		

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/misc_feature
13467. .13930
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fc94g02.y1"
13470. .13843
/notes="similar to Mus musculus EST AI060775 (NID:g3336198)"
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13728. .14181
/notes="similar to Danio rerio EST AW116140 (NID:g6082478)"
fi09f10.x1"
14189. .14189
/rpt_family="(CAAA)n"
14238. .14291
/rpt_family="(T)n"
14240. .14639
/notes="similar to Rattus norvegicus EST AA964818
(NID:g4279692)"
14249. .14403
/notes="similar to Rattus norvegicus EST AI704282
(NID:g4992182)"
14266. .14451
/notes="match to EST AA504508 (NID:g22406698) aa60g08.sl"
14287. .14414
/notes="similar to Danio rerio EST AW116140 (NID:g6082478)"
fi09f10.x1"
14405. .14886
/notes="similar to Rattus sp. EST AW143415 (NID:g6163418)"
14497. .14526
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14541. .14909
/notes="similar to Rattus norvegicus EST AI029866
(NID:g4301527)"

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Query Match	16.0%;	Score 97.2;	DB 86;	Length 152081;
Best Local Similarity	72.2%;	Pred. No. 1e-17;		
Matches 140;	Conservative 0;	Mismatches 53;	Indels 1;	Gaps 1;
17	ccccatccacgagtaattgtcaattccaaagacggtgttttgattgttcacacctgggt	76		
QY				
70146	CTCCACTCCCAAGGACATTGGCAATGCTAGAGACAT-TTTGGTTGTCACAACTCGGG	70088		
Db				
77	ggggaacatgctactggcatctaatgcatagagggcagtgtaatgctgctaaacatctttca	136		
QY				
70087	GGAGGGGGTGATACTGSCATCTAGTGATAGAGGCCGGGATGCTGTAAAAATCCTATA	70028		
Db				
137	acgcacagcagcagagcccccacaaagagaattatctagccccaaatgccataaacctgc	196		
QY				
70027	ATGCACAGGACAGACCCCCACCAAGAAATCTTCCTGCCCAAAATGTCACACAGTGCAGA	69968		
Db				
197	tgttgagaaaaacct	210		
QY				
69967	GGTTAAGAACCTCT	69954		
Db				

[illegible]

us-09-656-668-198.rge

Wed Nov 7 09:21:19 2001

CB10 ISA, UK. E-mail enquiries: humquerry@sanger.ac.uk Clone requests: clonerequests@sanger.ac.uk

On Mar 16, 2001 this sequence version replaced gi:10198608. During sequence assembly data is compared from overlapping clones where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em., EMBL; Sw., SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information on the WORMPEP database can be found at <http://www.sanger.ac.uk/projects/Celegans/wormpep> This sequence was generated from part of bacterial clone contigs of human chromosome 20, constructed by the Sanger Centre Chromosome 20 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr20>

RP5-998H6 is from the library RPCI-5 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>

VECTOR: pCYPAC2

This sequence is the entire insert of clone RP5-998H6 The true right end of clone RP4-599F21 is at 29571 in this sequence. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest.

Location/Qualifiers

1. .147815
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="20"
/map="q13.1"
/clone="RP5-998H6"
/clone_lib="RPCI-5"
85. .149
/note="MIR repeat: matches 60. .125 of consensus"
repeat_region
865. .905
/note="MER91 repeat: matches 153. .195 of consensus"
repeat_region
1141. .1358
/note="MIR repeat: matches 27. .250 of consensus"
repeat_region
1959. .2058
/note="L2 repeat: matches 2625. .2740 of consensus"
repeat_region
2081. .2176
/note="L2 repeat: matches 2612. .2710 of consensus"
repeat_region
2303. .2442
/note="7 copies 20 mer 81% conserved"
repeat_region
3585. .3790
/note="MIR repeat: matches 20. .239 of consensus"
repeat_region
4704. .4749
/note="23 copies 2 mer gt 84% conserved"
repeat_region
misc_feature
complement(4990. .5548)
/note="match: GSS: Em:AQ602943"
repeat_region
5125. .5354
/note="115 copies 2 mer tg 62% conserved"
repeat_region
5151. .5390
/note="12 copies 20 mer 60% conserved"
repeat_region
5397. .5442
/note="23 copies 2 mer gt 84% conserved"
repeat_region
5645. .5830
/note="MIR repeat: matches 1. .110 of consensus"
repeat_region
5831. .6141
/note="AluSq repeat: matches 1. .313 of consensus"
repeat_region
6142. .6394
/note="MIR repeat: matches 110. .390 of consensus"
repeat_region
6855. .7055
/note="MIR repeat: matches 59. .260 of consensus"
repeat_region
7129. .7544
/note="L2 repeat: matches 1529. .1989 of consensus"

7601. .8077
/note="MIR repeat: matches 34. .514 of consensus"
repeat_region
8089. .8174
/note="L2 repeat: matches 2066. .2155 of consensus"
repeat_region
8183. .8364
/note="MER53 repeat: matches 1. .182 of consensus"
repeat_region
10532. .10665
/note="MIR repeat: matches 80. .220 of consensus"
repeat_region
10708. .10764
/note="MIR repeat: matches 193. .251 of consensus"
repeat_region
11506. .11687
/note="MIR repeat: matches 29. .223 of consensus"
repeat_region
11702. .11810
/note="MIR repeat: matches 82. .190 of consensus"
repeat_region
11813. .12111
/note="Alu repeat: matches 1. .312 of consensus"
repeat_region
12113. .12306
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repeat_region
12123. .12302
/note="9 copies 20 mer 69% conserved"
repeat_region
12593. .12701
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repeat_region
13003. .13232
/note="L2 repeat: matches 2266. .2499 of consensus"
repeat_region
13372. .13455
/note="MIR repeat: matches 72. .161 of consensus"
repeat_region
13574. .13780
/note="MIR repeat: matches 56. .262 of consensus"
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/note="MIR repeat: matches 65. .262 of consensus"
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mRNA
complement(join(14281. .15619,18488. .18739,27130. .27247,27368. .27489,39965. .40101,50849. .51102,53537. .53730,57368. .57535,68050. .68169,81505. .81799,91582. .92237))
/gene="dJ998H6.1"
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/evidence="not_experimental"
/product="dJ998H6.1 (Ortholog of rat PB-Cadherin)"
gene
complement(14281. .92237)
/gene="dJ998H6.1"
polyA_signal
complement(14299. .14304)
misc_feature
14968. .15565
/note="CpG island"
/evidence="not_experimental"
complement(join(15048. .15619,18488. .18739,27130. .27247,27368. .27489,39965. .40101,50849. .51102,53537. .53730,57368. .57535,68050. .68169,81505. .81799,91582. .91836))
CDS
/gene="dJ998H6.1"
/note="match: proteins: Tr:Q63315 Tr:Q63561 Tr:Q90763 Tr:Q90762 Sw:P55285 Sw:Q13634 Sw:P79995 Tr:Q15066 Tr:Q90762 Sw:P55288 Sw:P55280 Sw:P97326 Tr:Q9JIM2 Sw:P55287 Sw:P52288 Tr:Q93319 Tr:Q9ZOM3 Tr:Q9WTP5 Tr:O43205 Tr:Q9ULB2 Tr:P55286 Tr:O54800 Sw:P97291 Sw:P55289 Tr:Q9ULB5 Tr:Q9ULB4"
/codon_start=1
/evidence="not_experimental"
/product="dJ998H6.1 (ortholog of rat PB-Cadherin)"
/protein_id="CAB51587.2"
/db_xref="GI:13366277"
/db_xref="SPTREMBL:Q9UL99"
/translation="MRPRGRLRAGVALSPALLILLPPPTLLGLRLAAGTPSP SAGARQDAGLGGKRWGWNQFFVEEYTGTEPLYVCKIHSDEGDAIKYIS GEAGATFLIDELTGDIHAMELRDQKFTYTLRAQDRATNRLLEPESEFLIKVQ INDESPRFLHGPYIGSVAELSPGTSQVMQMASDADPTYGSSARLYSVLDGHHFT VDFKGTGIVTAVPDLDRSQREVEVVTQATDMAGQLGSGSTVTIIVTVDNDNPPR FPQKMQYQIOESAPITAGVGRKKAEDSDVGENTDMTYHLKDESSGGGVFKVTTDSD

REFERENCE	AUTHORS	JOURNAL	TITLE	COMMENT
1 (bases 1 to 185143)	Birten, B., Linton, L., Nusbaum, C. and Lander, E.	Unpublished	Homo sapiens, clone RP11-766D10	
2 (bases 1 to 185143)	Birten, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N., Anderson, S., Baldwin, J., Barna, N., Bastien, V., Beda, F., Boquslavsky, L., Boukhgalter, B., Brown, A., Burkett, G., Campomiano, A., Cooke, P., DeAtrellano, K., Dewar, K., Diaz, J.S., Collymore, A., Cooke, P., DeAtrellano, K., Dewar, K., Diaz, J.S., Dodge, S., Domino, M., Doyle, M., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N., Grant, G., Hagos, B., Jones, C., Kann, L., Karatas, A., Howland, J.C., Iliev, I., Johnson, R., Landers, T., Lenockzy, J., Levine, R., Liu, C., Liu, G., Locke, K., Macdonald, P., Marquis, N., McCarthy, M., McEwan, P., McGurk, A., McKernan, K., McPheeters, R., Melchior, J., Meneus, L., Mihova, T., Miranda, C., Mlenka, V., Morrow, J., Murphy, T., Naylor, J., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neil, D., Olivari, T.M., Oliver, J., Peterson, K., Pierre, N., Pisani, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rothman, D., Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J., Tirrell, A., Travers, M., Trigilio, J., Vassiliou, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G., Zainoun, J., Zimmer, A. and Zody, N.	Whitehead Institute/MIT Center for Genome Research	Submitted (16-MAR-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	On Aug 26, 2000 this sequence version replaced gi:9887743. All repeats were identified using RepeatMasker: Smit, A.F.A. & Green, P. (1996-1997) http://ftp.genome.washington.edu/RM/RepeatMasker.html
Center code: W1BR	Center site: http://www-seq.wi.mit.edu	Contact: sequence_submissions@genome.wi.mit.edu	Project Information	Center project name: L7554
Center clone name: 766_D_10	Summary Statistics	Sequencing vector: M13: M77815; 100% of reads	Chemistry: Dye-terminator Big Dye; 100% of reads	Assembly program: Phrap; version 0.960731
Consensus quality: 174463 bases at least Q30	Consensus quality: 178842 bases at least Q20	Insert size: 200000; agarose-fp	Insert size: 180843; sum-of-contigs	Quality coverage: 2.9 in Q20 bases; agarose-fp
Quality coverage: 3.2 in Q20 bases; sum-of-contigs	NOTE: This is a 'working draft' sequence. It currently consists of 44 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.	1 200: contig of 200 bp in length	201 300: gap of 100 bp	301 1640: contig of 1340 bp in length
1641 1740: gap of 100 bp	1741 3026: contig of 1286 bp in length	3027 3126: gap of 100 bp	3127 4675: contig of 1549 bp in length	4676 4775: gap of 100 bp
4776 5875: contig of 1000 bp in length	5876 5875: gap of 100 bp	5876 7301: contig of 1426 bp in length	7301 7401: gap of 100 bp	

* 170175 170274: gap of 100 bp
170275 170374: gap of 14869 bp in length.

*	7402	8749:	contig of 1348 bp in length
*	7500	8849:	gap of 100 bp
*	8850	10630:	contig of 1781 bp in length
*	10631	10730:	contig of 100 bp
*	10731	12287:	contig of 1557 bp in length
*	12288	12367:	gap of 100 bp
*	12388	14291:	contig of 1904 bp in length
*	14292	14391:	gap of 100 bp
*	14392	15981:	contig of 1590 bp in length
*	15982	16081:	gap of 100 bp
*	16082	17373:	contig of 1292 bp in length
*	17374	17473:	gap of 100 bp
*	17474	19023:	contig of 1550 bp in length
*	19024	19123:	gap of 100 bp
*	19124	20568:	contig of 1445 bp in length
*	20569	20668:	gap of 100 bp
*	20669	22533:	contig of 1865 bp in length
*	22534	22633:	gap of 100 bp
*	22634	24930:	contig of 2297 bp in length
*	24931	25030:	gap of 100 bp
*	25031	27214:	contig of 2184 bp in length
*	27215	27314:	gap of 100 bp
*	27315	30058:	contig of 2744 bp in length
*	30059	30158:	gap of 100 bp
*	30159	31989:	contig of 1831 bp in length
*	31990	32089:	gap of 100 bp
*	32090	35144:	contig of 3055 bp in length
*	35145	35244:	gap of 100 bp
*	35245	37733:	contig of 2489 bp in length
*	37734	37833:	gap of 100 bp
*	37834	40275:	contig of 2442 bp in length
*	40276	40375:	gap of 100 bp
*	40376	42771:	contig of 2396 bp in length
*	42772	42871:	gap of 100 bp
*	42872	46769:	contig of 3898 bp in length
*	46770	46869:	gap of 100 bp
*	46870	49648:	contig of 2779 bp in length
*	49649	49748:	gap of 100 bp
*	49749	53828:	contig of 4080 bp in length
*	53829	53928:	gap of 100 bp
*	53929	58678:	contig of 4750 bp in length
*	58679	58778:	gap of 100 bp
*	58779	63068:	contig of 4290 bp in length
*	63069	63168:	gap of 100 bp
*	63169	67733:	contig of 4565 bp in length
*	67734	67833:	gap of 100 bp
*	67834	72886:	contig of 5053 bp in length
*	72887	72986:	gap of 100 bp
*	72987	76848:	contig of 3862 bp in length
*	76849	76948:	gap of 100 bp
*	76949	82756:	contig of 5808 bp in length
*	82757	82856:	gap of 100 bp
*	82857	88725:	contig of 5869 bp in length
*	88826	88825:	gap of 100 bp
*	88826	95518:	contig of 6693 bp in length
*	95519	95618:	gap of 100 bp
*	95619	101325:	contig of 5707 bp in length
*	101326	101425:	gap of 100 bp
*	101426	106203:	contig of 4778 bp in length
*	106204	106303:	gap of 100 bp
*	106304	111060:	contig of 4757 bp in length
*	111061	111160:	gap of 100 bp
*	111161	119653:	contig of 8493 bp in length
*	119654	119753:	gap of 100 bp
*	119754	127901:	contig of 8148 bp in length
*	127902	128001:	gap of 100 bp
*	128002	136682:	contig of 8681 bp in length
*	136683	136782:	gap of 100 bp
*	136783	146372:	contig of 9590 bp in length
*	146373	146472:	gap of 100 bp
*	146473	157184:	contig of 10712 bp in length
*	157185	157284:	gap of 100 bp
*	157285	170174:	contig of 12890 bp in length

FEATURES	source	Location/Qualifiers
		1. .185143
		/organism="Homo sapiens"
		/db_xref="taxon:9606"
		/clone_xref="RP11-766D10"
		/clone_lib="RPC1-11 Human Male BAC"
misc_feature		1. .200
		/note="assembly_fragment"
		clone_end:SP6
		vector_side:left"
		301. .1640
misc_feature		/note="assembly_fragment"
misc_feature		1741. .3026
		/note="assembly_fragment"
misc_feature		3127. .4675
		/note="assembly_fragment"
misc_feature		4776. .5775
		/note="assembly_fragment"
misc_feature		5876. .7301
		/note="assembly_fragment"
misc_feature		7402. .8749
		/note="assembly_fragment"
misc_feature		8850. .10630
		/note="assembly_fragment"
misc_feature		10731. .12287
		/note="assembly_fragment"
misc_feature		12388. .14291
		/note="assembly_fragment"
misc_feature		14392. .15981
		/note="assembly_fragment"
misc_feature		16082. .17373
		/note="assembly_fragment"
misc_feature		17474. .19023
		/note="assembly_fragment"
misc_feature		19124. .20568
		/note="assembly_fragment"
misc_feature		20669. .22533
		/note="assembly_fragment"
misc_feature		22634. .24930
		/note="assembly_fragment"
misc_feature		25031. .27214
		/note="assembly_fragment"

Query Match	15.4%;	Score 93.6;	DB 69;	Length 185143;
Best Local Similarity	66.2%;	Pred. No. 1.2e-16;		
Mismatches 115;	Conservative	0;	Mismatches 69;	Indels 0;
Gaps 0;				

[illegible]

RESULT	9
AL353791/c	DNA
LOCUS	AL353791 208763 bp HTG
DEFINITION	Homo sapiens chromosome 9 clone RP11-475B17, *** SEQUENCING INFORMATION ***
ACCESSION	AL353791
VERSION	AL353791.5 GI:13373919 PROGRESS ***, 4 unordered pieces.
	12-MAR-2001

[illegible]

RESULT	10			PRI	05-MAY-2000
AC016951			DNA		
LOCUS	Homo sapiens	3 BAC RP11-452H12	(Roswell Park Cancer Institute Human		
DEFINITION	BAC library)	complete sequence.			
ACCESSION	AC016951				
VERSION	AC016951.9	GI:7658300			
KEYWORDS	HTG.				
SOURCE	human.				
ORGANISM	Homo sapiens				
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
	Mammalia; Primates; Hominoidea; Hominidae; Homo.				

1 (bases 1 to 147929)

REFERENCE
AUTHORS

Muzny D.M., Adams, C., Bailey, M., Barbara, J., Blankenburg, K.,
Bodota, B., Bouck, J., Bowie, S., Brooks, A., Bunay, C., Bunac, C.,
Burkett, C., Burrows, J., Carter, M., Chacko, J., Chen, Z., Cox, C.,
David, K., Delgado, O., Deshazo, D., Ding, Y., Domah-Rasthid, N.,
Dugan-Kocha, S., Durbin, K.J., Fernandez, C., Ferraguto, D.,
Forcum-Tansey, J., Frantz, P., Ganesh, R., Garcia, D.K., Correll, J.H.,
Gorell, L.L., Guevara, W., Harris, K., He, X., Hernandez, J.,
Hodgson, A., Hoques, M., Holloway, C., Hosak, K., Jackson, L.E.,
Jackson, L., Jia, Y., Jones, M., Kelly, S., Kondjewski, N., Kong, Y.,
Kovar, C., Leal, P., Li, Z., Lichtarge, O., Liu, J., Liu, W., Logan, O.P.,
Lozado, R.J., Lu, J., Lucier, R., Martin, R., Martinez, C., Nash, S.,
Mei, G., Moore, S., Moorish, I., Morgan, M., Morris, S., Oswal, G., Parish, B.,
Nelson, A., Nguyen, R., Nguyen, N., Nguyen, S., Quiles, E., Shen, H.,
Paxton, S., Payton, B., Perez, L., Pu, L., Quiles, M., Reiter, D.,
Rives, M., Samuel, S., Say, J., Scherret, S., Sugang, R., Tabor, P., Taylor, T.,
Simon, M., Sparks, A., Stamps, A., Sucgang, R., Watlington, S.,
Vasquez, L., Vinson, R., Vo, Q., Wahbah, M., Worley, K., Wren, J.,
Weinstock, G., Weinstock, I.R., Williamson, A., Wyden, K., and Gibbs, R.

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

Direct Submission
Unpublished
2 (bases 1 to 147929)
Worley, K.C.
Direct Submission
Submitted (09-DEC-1999) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
TX 77030, USA

REFERENCE 3 (bases 1 to 147929)
AUTHORS Worley, K.C.
TITLE Direct Submission
JOURNAL Submitted (28-APR-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
REFERENCE 4 (bases 1 to 147929)
AUTHORS Worley, K.C.

JOURNAL
TITLE Submitted (05-MAY-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
COMMENT On Apr 28, 2000 this sequence version replaced gi:7331304.
Information: <http://www.hgsc.bcm.tmc.edu/> or email acsb@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent a

KEYWORDS	HTG: HTGS_PHASE1; HTGS_DRAFT.
SOURCE	human.
ORGANISM	Homo sapiens
REFERENCE	Craniata: Vertebrata; Euteleostomi; Mamalia: Eutheria; Primates; Homo. 1 (bases 1 to 208763)
AUTHORS	Plumb, B.
TITLE	Direct Submission
JOURNAL	Submitted (11-Mar-2001) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk requests: clonerquest@sanger.ac.uk
COMMENT	On Mar 16, 2001 this sequence version replaced gi:11340258. On Mar 16, 2001 this sequence version replaced gi:11340258

```

Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk
----- Project Information -----
Center project name: BA472R14
----- Summary Statistics -----
Assembly program: XGAP4; version 4.5
Sequencing vector: plasmid; L08752; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Consensus quality: 208161 bases at least Q40
Consensus quality: 208281 bases at least Q30
Consensus quality: 208368 bases at least Q20
Insert size: 208463; sum-of-contigs
Insert size: 209014; 4.8% error; agarose-fp
Quality coverage: 11.65x in Q20 bases; sum-of-contigs
Quality coverage: 11.61x in Q20 bases; agarose-fp

```

* NOTE: This is a 'working draft' sequence. It currently
* consists of 4 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

*	i	119577: contig of 119577 bp in length
*		119578 119677: gap of 100 bp
*		119678 144219: contig of 24542 bp in length
*		144220 144319: gap of 100 bp
*		144320 204287: contig of 59968 bp in length
*		204288 204387: gap of 100 bp
*		204388 208753: contig of 4376 bp in length.

FEATURES	Location/Qualifiers
source	1. .208763
	/organism="Homo sapiens"
	/db_xref="taxon:9606"
	/chromosome="9"
	/clone="rp11-475B17"
	/clone_lib="RPC1-11.2"
misc_feature	1. .119577
	/note="assembly_fragment:00460
	fragment_chain:1"
misc_feature	119678. .144219
	/note="assembly_fragment:00991
	fragment_chain:1"
misc_feature	144320. .204287
	/note="assembly_fragment:04392
	fragment_chain:1"
misc_feature	204388. .208763
	/note="assembly_fragment:05279.0"
BASE COUNT	60862 a 42636 c 42179 g 62780 t
ORIGIN	306 others

Query Match 15.4%; Score 93.6; DB 79; Length 208763;
Best Local Similarity 66.2%; Pred. No. 1.2e-16; Models 0; Gaps 0;

tttctccacacctatattgattatcacaccctgggtcgggaacatgccta 89

entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:
STSs are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.
Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished) for Human and Mouse sequences.
Genes and region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://gc.bcm.tmc.edu:8088/quality/info/genbank.annotation.html>.

QUALSTAT-REPORT-----

----- Summary Statistics -----
Contig length: 147929
Phrap values in estimate: 146381
Average error rate (BCM-Phrap estimate): 0.000137522
Fraction of Phrap values less than 40 : 0.0255019
Number of consensus changing edits: 17
Number of N's in consensus : 0

----- Consensus changing edits -----
Position Original+Context Edited+Context
26320 acctaatgct(n)gatgacgagt acctaatgct(a)gatgacgagt
31259 ttcttttgg(n)aagatgaac ttcttttgg(g)aagatgaac
37952 gaataatt(n)gaataaagat gaataatt(g)gaataaagat
48020 agattcataa(n)gcaagtcctg agattcataa(a)gcaagtcctg
61999 ttctaaaatt(n)gttttgactt ttctaaaatt(a)gttttgactt
72566 tgattttgtt(n)cttgggaagc tgattttgtt(t)cttgggaagc
81082 ctcatatagc(n)tacatactag ctcatatagc(c)tacatactag
81162 atgaagtaag(n)caagccatg atgaagtaag(a)caagccatg
81177 gccatggagg(n)agatggagg gccatggagg(g)agatggagg
82666 aatttggaaa(n)aatccacatc aatttggaaa(c)aatccacatc
91636 atgatagggt(n)tacacaaatc atgatagggt(t)tacacaaatc
91636 cgtccatgcn(c)caaaaaaaa cgtccatgcn(t)caaaaaaaa
96541 aaaaaaaatt(t)aaaaaaaaa aaaaaaaatt(a)aaaaaaaaa
96541 aaaaaaaag(g)aaggtgtca aaaaaaaag(g)aaggtgtca
98037 gctgcaagca(n)gccgggaagc gctgcaagca(c)gccgggaagc
139118 gaggcagagg(n)caaggaaagg gaggcagagg(c)caaggaaagg

----- Distribution of Quality < 40 Bases -----

1000	*	*
900	*	*
800	*	*
700	*	*
#	*	*
bases 600		

FEATURES	source	Version: 1.01	qxfo.	Location/Qualifiers	Phrap Value Range
repeat_region	STS	1.147929		/organism="Homo sapiens"	5 10 15 20 25 30 35 40
repeat_region	STS			/db_xref="taxon:9606"	
repeat_region	STS			/chromosome="3"	
repeat_region	STS			/clone="RP11-452H12"	
repeat_region	STS			/complement(532..617)	
repeat_region	STS			/rpt_family="MSTB"	
repeat_region	STS			625..696	
repeat_region	STS			/rpt_family="LTR9"	
repeat_region	STS			668..891	
repeat_region	STS			/standard_name="Z53523"	
repeat_region	STS			/db_xref="dbSTS:45636"	
repeat_region	STS			697..731	
repeat_region	STS			/rpt_family="(CA)n"	
repeat_region	STS			complement(732..1090)	
repeat_region	STS			/rpt_family="MSTB"	
repeat_region	STS			1927..2384	
repeat_region	STS			/rpt_family="MLT1G1"	
repeat_region	STS			3079..3391	
repeat_region	STS			/rpt_family="AluJb"	
repeat_region	STS			complement(3685..3897)	
repeat_region	STS			/rpt_family="MER20"	
repeat_region	STS			complement(5392..5890)	
repeat_region	STS			/rpt_family="Charliea"	
repeat_region	STS			complement(5935..6618)	
repeat_region	STS			/rpt_family="Charliea"	
repeat_region	STS			8265..8469	
repeat_region	STS			/standard_name="G09350"	
repeat_region	STS			/db_xref="dbSTS:15731"	
repeat_region	STS			8348..8390	
repeat_region	STS			/rpt_family="(TTA)n"	
repeat_region	STS			8503..8814	
repeat_region	STS			/rpt_family="AluSx"	
repeat_region	STS			9081..9226	
repeat_region	STS			/rpt_family="MER53"	
repeat_region	STS			9245..9413	
repeat_region	STS			/rpt_family="MSTD"	
repeat_region	STS			9414..9705	
repeat_region	STS			/rpt_family="AluJb"	
repeat_region	STS			9706..9951	
repeat_region	STS			/rpt_family="MSTD"	
repeat_region	STS			9952..10035	
repeat_region	STS			/rpt_family="MER53"	
repeat_region	STS			11657..13087	
repeat_region	STS			/rpt_family="pTR5"	
repeat_region	STS			complement(13237..13437)	
repeat_region	STS			/rpt_family="MLT1H"	
repeat_region	STS			19462..19885	
repeat_region	STS			/rpt_family="HAL1"	
repeat_region	STS			20035..20336	
repeat_region	STS			/rpt_family="AluJb"	
repeat_region	STS			20395..21174	
repeat_region	STS			/rpt_family="L1PA2"	
repeat_region	STS			21170..26420	
repeat_region	STS			/rpt_family="L1PA2"	
repeat_region	STS			25950..26082	
repeat_region	STS			/standard_name="G19948"	
repeat_region	STS			/db_xref="dbSTS:32826"	

[illegible]

Query Match	15.1%	Score 91.8:	DB 87:	Length 147929;
Best Local Similarity	64.2%;	Pred. No. 4e-16;		
Matches 138; Conservative	0;	Mismatches 77;	Indels 0;	Gaps 0;
2	gagttgccccctaccccccatccccagtgaaatttcgaattcctaagaagcagtgatttga	61		
28127	GAATTTGGCACCACCCACCCAGGAATAATTAAACAATTTCTCGAGATATTTTGGC	28186		
62	tgtcacacctgggtggggaaca tgc tactggc atcta atgc atag aggc agta atgct	121		
28187	TGTCACACTTGAGGAGAGGGGTGCTACTGGCATCTAATGAATGAGGCCAAGATCCT	28246		
122	gcttaacatcttttaacgcacagagacagagcccccacaaagagaattatctagccccaaa	181		
28247	GCTAAATCTTCTACAATGCACAGGACAGCTCCCCCAACACAAAGAATTATTTCAGACCAAA	28306		
182	tgccaataacactgctgttgagaaaaacctaccgcga	216		
28307	TTTCACATAATGCAGAGTGCAGAAAACACTGCTGTA	28341		

RESULT	11	AL353729	194313 bp	DNA	HTG	06-MAR-2001
LOCUS		Homo sapiens chromosome 9 clone RP11-290L7, *** SEQUENCING IN				
DEFINITION		Homo sapiens chromosome 9 clone RP11-290L7, *** SEQUENCING IN				
PROGRESS		Homo sapiens chromosome 9 clone RP11-290L7, *** SEQUENCING IN				
ACCESSION		AL353729				
VERSION		AL353729.7	GI:13273646			
KEYWORDS		HTG: HTGS_PHASE1; HTGS_DRAFT.				
SOURCE		human.				
ORGANISM		Homo sapiens				
REFERENCE		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
AUTHORS		Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
TITLE		1 (bases 1 to 194313)				
JOURNAL		Plumb,B.				
COMMENT		Direct Submission				
		Submitted (03-MAR-2001) Sanger Centre, Hinxton, Cambridgeshire,				
		CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk				
		requests: clonerequest@sanger.ac.uk				
		On Mar 11, 2001 this sequence version replaced gi:9800794.				
		----- Genome Center				
		Center: Sanger Centre				
		Center code: SC				
		Web site: http://www.sanger.ac.uk				
		Contact: humquery@sanger.ac.uk				
		----- Project Information				
		Center project name: BA290L7				
		Center project name: BA290L7				

```

----- Summary Statistics -----
Assembly program: XGAP4; version 4.5
Sequencing vector: plasmid; L08752; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Consensus quality: 193402 bases at least Q40
Consensus quality: 193962 bases at least Q30
Consensus quality: 194209 bases at least Q20
Insert size: 194313; sum-of-contigs
Insert size: 196358; 3.1% error; agarose-fp
Quality coverage: 6.31x in Q20 bases; sum-of-contigs Quality
coverage: 6.45x in Q20 bases; agarose-fp
-----
* NOTE: This is a 'working draft' sequence.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

FEATURES
    source                Location/Qualifiers
        1..194313
            /organism="Homo sapiens"
            /db_xref="taxon:9606"
            /chromosome="9"
            /clone="RP11-290L7"
            /clone_lib="RP11-11.2"
        1..194313
            /note="assembly_fragment:01008
clone end:SP6
misc_feature

```

```
misc_feature
1. .194313
/note="assembly_fragment:01008
clone_end:SP6
```


* arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

1 3206: contig of 3206 bp in length
 * 3207: gap of unknown length
 * 3307: contig of 8451 bp in length
 * 11757: gap of unknown length
 * 11758: contig of 13077 bp in length
 * 24934: gap of unknown length
 * 25034: contig of 11537 bp in length
 * 25035: gap of unknown length
 * 36571: gap of unknown length
 * 36572: contig of 14049 bp in length
 * 50720: gap of unknown length
 * 50721: contig of 10844 bp in length
 * 50821: gap of unknown length
 * 61664: gap of unknown length
 * 61665: contig of 15476 bp in length
 * 61765: gap of unknown length
 * 77241: gap of unknown length
 * 77242: contig of 27656 bp in length
 * 104997: gap of unknown length
 * 105096: contig of 27667 bp in length
 * 132763: gap of unknown length
 * 132764: contig of 27317 bp in length
 * 132864: gap of unknown length
 * 160180: contig of 4642 bp in length
 * 160281: gap of unknown length
 * 164923: contig of unknown length
 * 165023: gap of unknown length
 * 166928: contig of 1905 bp in length
 * 167028: gap of unknown length
 * 170224: contig of 3197 bp in length
 * 170225: gap of unknown length
 * 173939: contig of 3615 bp in length
 * 173940: gap of unknown length
 * 174040: contig of 7062 bp in length
 * 181101: gap of unknown length
 * 181201: contig of 8178 bp in length
 * 181202: gap of unknown length
 * 189380: contig of 9706 bp in length
 * 189480: gap of unknown length
 * 199186: contig of 8461 bp in length.
 * 199286: Location/Qualifiers

FEATURES

source
 1. 207746
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="UL"
 /clone="RP11-566J10"
 1. 3206
 /note="assembly_name:Contig1"
 3307. 11757
 /note="assembly_name:Contig10"
 11858. 24934
 /note="assembly_name:Contig11"
 25035. 36571
 /note="assembly_name:Contig12"
 clone_end:T7
 vector_side:right
 36672. 50720
 /note="assembly_name:Contig13"
 50821. 61664
 /note="assembly_name:Contig14"
 clone_end:SP6
 vector_side:left
 61765. 77240
 /note="assembly_name:Contig15"
 77341. 104996
 /note="assembly_name:Contig16"
 105097. 132763
 /note="assembly_name:Contig17"
 132864. 160180
 /note="assembly_name:Contig18"
 160281. 164922
 /note="assembly_name:Contig19"
 165023. 166927

/note="assembly_name:Contig3"
 167028. 170224
 /note="assembly_name:Contig4"
 170325. 173939
 /note="assembly_name:Contig5"
 174040. 181101
 /note="assembly_name:Contig6"
 181202. 189379
 /note="assembly_name:Contig7"
 189480. 199185
 /note="assembly_name:Contig8"
 199286. 207746
 /note="assembly_name:Contig9"
 60979 a 43162 c 40763 g 61116 t 1726 others
 BASE COUNT
 ORIGIN

Query Match 15.0%; Score 91; DB 64; Length 207746;
 Best Local Similarity 66.3%; Pred. No. 7.1e-16;
 Matches 130; Conservative 0; Mismatches 66; Indels 0; Gaps 0;
 QY 24 cccagtgaaatttgcaattcctaaagacgtgtttttgattgtcacaccctgggtgggaac 83
 Db 87414 CCCAGGACATGTGGCAATGTTTGAGACACTTTAGTGTGTCAAACTTGGGGAGGTG 87355
 QY 84 atgtactggcatctaatgcatagaggcagtaagtctgctaaacattcttcaacgcaca 143
 Db 87354 GTGCTAGTGGCAATTTAGTGTAGGTAGAGGTGACAGACACTGCTAAACATCTCTGCAGACTGAG 87295
 QY 144 ggcagagcccccaaaagagaatattctagcccccaaatgtccataacactgtgtgag 203
 Db 87294 GGACGACACCCACACAAAGAAATATCTCTCTAAATGACAAATGTGCTGAGACTGAG 87235
 QY 204 aaacactaccgcagga 219
 Db 87234 MAGCCTGCTTTAGGA 87219

RESULT 13
 AC021051 162082 bp DNA HTG 04-NOV-2000
 LOCUS Homo sapiens chromosome 3 clone RP11-214M2, WORKING DRAFT SEQUENCE,
 DEFINITION 6 unordered pieces.

ACCESSION AC021051
 VERSION AC021051.9 GI:11079339
 KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
 SOURCE human.

ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Cranialata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 162082)

REFERENCE
 AUTHORS
 Alsbrooks S.L., Amaralunge H.C., Are J.R., Banks T., Barbacia J.,
 Benton J., Bimaga K., Blankenburg K., Bonnin D., Bouck J.,
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* * * 13060 13753: contig of 694 bp in length
* * * 13754 13853: gap of 100 bp
* * * 13854 14552: contig of 699 bp in length
* * * 14552 14652: gap of 100 bp
* * * 14553 15362: contig of 710 bp in length
* * * 15363 15462: gap of 100 bp
* * * 15463 16265: contig of 703 bp in length
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* * * 16969 17068: gap of 100 bp
* * * 17069 17792: contig of 724 bp in length
* * * 17793 17892: gap of 100 bp
* * * 17893 18601: contig of 709 bp in length
* * * 18602 18701: gap of 100 bp
* * * 18702 19397: contig of 696 bp in length
* * * 19398 19497: gap of 100 bp
* * * 19498 20203: contig of 706 bp in length
* * * 20204 20303: gap of 100 bp
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* * * 20995 21094: gap of 100 bp
* * * 21095 21795: contig of 701 bp in length
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* * * 21896 22600: contig of 705 bp in length
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* * * 24287 24999: contig of 713 bp in length
* * * 25000 25099: gap of 100 bp
* * * 25100 25860: contig of 761 bp in length
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* * * 41372 42084: contig of 713 bp in length
* * * 42085 42184: gap of 100 bp
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 Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
 Tirrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J.,
 Zimmer,A. and Zody,M.
 Direct Submission
 Submitted (06-FEB-2000) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Mar 18, 2001 this sequence version replaced gi:9127546.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
 ----- Genome Center / MIT Center for Genome Research
 Center code: WIBR
 Web site: <http://www.seq.wi.mit.edu>
 Contact: sequence_submissions@genome.wi.mit.edu
 ----- Project Information
 Center project name: 14317
 Center clone name: 659_D_6

 * NOTE: This record contains 78 individual
 * sequencing reads that have not been assembled into
 * contigs. Runs of N are used to separate the reads
 * and the order in which they appear is completely
 * arbitrary. Low-pass sequence sampling is useful for
 * identifying clones that may be gene-rich and allows
 * overlap relationships among clones to be deduced.
 * However, it should not be assumed that this clone
 * will be sequenced to completion. In the event that
 * the record is updated, the accession number will
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 * 1 705: contig of 705 bp in length
 * 706 805: gap of 100 bp
 * 806 1565: contig of 760 bp in length
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 * 3186 3285: gap of 100 bp
 * 3286 4007: contig of 722 bp in length
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 * 7281 7380: gap of 100 bp
 * 7381 8087: contig of 707 bp in length
 * 8088 8187: gap of 100 bp
 * 8188 8895: contig of 708 bp in length
 * 8896 8995: gap of 100 bp
 * 8996 9724: contig of 729 bp in length
 * 9725 9824: gap of 100 bp
 * 9825 10558: contig of 734 bp in length
 * 10559 10658: gap of 100 bp
 * 10659 11337: contig of 679 bp in length
 * 11338 11437: gap of 100 bp
 * 11438 12151: contig of 714 bp in length
 * 12152 12251: gap of 100 bp
 * 12252 12959: contig of 708 bp in length
 * 12960 13059: gap of 100 bp

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* 46987 47700: contig of 714 bp in length
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QY 84 atgtactggcatctaatgcatagagggcagtaatctgctaacaatctttcaacgcaca 143
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Search completed: November 5, 2001, 23:02:50
Job time: 2800 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model
Run on: November 5, 2001, 22:57:50 ; Search time 1188.78 Seconds
(without alignments)
7884.937 Million cell updates/sec

Title: US-09-656-668-198
Perfect score: 606
Sequence: 1 tgagtgtgccccctaccccc.....aagcctgtctgtctgcac 606

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0
Searched: 1344157 seqs, 7733874588 residues

Word size : 0
Total number of hits satisfying chosen parameters: 2688314

Minimum DB seq length: 0
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Post-processing: Listing first 45 summaries

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97:	gb_pr10:*
98:	em_ba3:*

No. is the number
greater than or equal
derived by analysis

Score	Query Match %	Length	ID	
506	100.0	606	AX093380	
59	9.7	90497	HS732E4	
26	4.3	105137	HSJ104017	
25	4.1	154969	AC012443	
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25	4.1	194294	AC079344	
7	23	3.8	67084	78
23	3.8	93959	92	

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

9	23	3.8 103757	75	AC073940	Homo sapi
10	23	3.8 127590	85	AC002554	Human Chr
11	23	3.8 134394	87	AC067916	Homo sapi
12	23	3.8 151357	62	AC015713	Homo sapi
13	23	3.8 157402	61	AC034161	Homo sapi
14	23	3.8 159723	66	AC021862	Homo sapi
15	23	3.8 171589	75	AC078953	Homo sapi
16	23	3.8 172214	64	AC016155	Homo sapi
17	23	3.8 172588	60	AC008590	Homo sapi
18	23	3.8 173808	71	AC036206	Homo sapi
19	23	3.8 174128	89	AC036206	Human DNA
20	23	3.8 176092	79	AC068222	Homo sapi
21	23	3.8 178028	79	AL355338	Homo sapi
22	23	3.8 178328	65	AC019265	Homo sapi
23	23	3.8 180316	86	AC007773	Homo sapi
24	23	3.8 186292	72	AC064876	Homo sapi
25	23	3.8 186699	77	AC030015	Homo sapi
26	23	3.8 189079	72	AC053545	Homo sapi
27	23	3.8 190009	68	AC090519	Homo sapi
28	23	3.8 191060	76	AC021468	Homo sapi
29	23	3.8 194937	68	AC024534	Homo sapi
30	23	3.8 207221	82	AC012564	Homo sapi
31	23	3.6 161837	86	AC006479	Homo sapi
32	22	3.6 174380	64	AC021607	Homo sapi
33	22	3.6 178042	68	AC023986	Homo sapi
34	22	3.6 194058	86	AC007554	Homo sapi
35	22	3.6 225635	80	AC007553	Homo sapi
36	21	3.5 39540	85	AC000079	Homo sapi
37	21	3.5 43461	6	CER0307	Z46828 Caenorhabdi
38	21	3.5 43934	85	AC000068	Homo sapi
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41	21	3.5 137658	92	HS1022J11	Human DNA
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44	21	3.5 150803	81	AL354985	Human DNA
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ALIGNMENTS

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RESULT 1
AX093380 606 bp DNA PAT 30-MAR-2001
LOCUS
SEQUENCE 198 from Patent WO0118046.
AX093380
ACCESSION
AX093380.1 GI:13509828
KEYWORDS
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 606)
XU,J. and Stolk,J.A.
AUTHORS
Ovarian tumor sequences and methods of use therefor
TITLE
Patent: WO 0118046-A 198 15-MAR-2001;
JOURNAL
CORIXA CORPORATION (US)
FEATURES
Location/Qualifiers
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/db_xref="taxon:9606"
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BASE COUNT
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Best Local Similarity 100.0%; Pred. No. 0; Gaps
Matches 606; Conservative 0; Mismatches 0; Indels 0;
QY 1 tgagttgccccctaccgccatcccgatgaattattgcgaattcctaaagaogtgttg 60
db 1 TGAGTTGGCCCTTACCCCAATCCAGTGAATTTGCAATTCCTAAACACCGTGTG 60

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together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information from the WORMPEP database can be found at <http://www.sanger.ac.uk/Projects/C.elegans/wormpep> CTA-732E4 is from the human BAC library described in U-J. Kim et al. (1996) Genomics 34, 213-218.

VECTOR: pBeloBAC11

This sequence is the entire insert of clone CTA-732E4 The true left end of clone RP11-541016 is at 5510 in this sequence. The true right end of clone CTA-544A11 is at 41939 in this sequence.

FEATURES

source

1. 90497
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 /db_xref="taxon.9606"
 /chromosome="22"
 /map="q12.1"
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 repeat_region 1374. 1869
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 repeat_region 3423. 3478
 /note="14 copies 4 mer caca 94 conserved"
 repeat_region 3670. 3742
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 repeat_region 3763. 4128
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 repeat_region 4612. 4930
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 repeat_region 4934. 5220
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 repeat_region 26003. 26192
 /note="MIR repeat: matches 20. .216 of consensus"
 repeat_region 27067. 27138
 /note="Charlie4 repeat: matches 1902. .1961 of consensus"
 repeat_region 27363. 27661
 /note="AluYb8 repeat: matches 1. .306 of consensus"
 repeat_region 28662. 28722
 /note="I2 repeat: matches 2683. .2744 of consensus"
 repeat_region 28758. 28797
 /note="10 copies 4 mer tata 87 conserved"
 repeat_region 28759. 28798
 /note="20 copies 2 mer at 87 conserved"
 repeat_region 29334. 29421
 /note="22 copies 4 mer gaaa 65 conserved"
 repeat_region 29805. 30155
 /note="L1MB5 repeat: matches 4639. .4984 of consensus"
 repeat_region 30150. 31160
 /note="L1MB5 repeat: matches 5157. .6166 of consensus"
 repeat_region 31294. 31449
 /note="L1PA2 repeat: matches 5991. .6146 of consensus"
 repeat_region 31544. 31681
 /note="MIR repeat: matches 93. .254 of consensus"
 repeat_region 31914. 32062
 /note="MIR repeat: matches 101. .261 of consensus"
 repeat_region 32553. 32814
 /note="I2 repeat: matches 1072. .1341 of consensus"
 repeat_region 32884. 32952
 /note="MER45B repeat: matches 2. .76 of consensus"
 repeat_region 33181. 33436
 /note="MER45C repeat: matches 676. .952 of consensus"
 repeat_region 33504. 33671
 /note="MIR repeat: matches 4. .191 of consensus"
 repeat_region 33771. 34072
 /note="AluJo repeat: matches 1. .310 of consensus"
 misc_feature complement(35126. 35616)
 /note="match: GSS: Em:AQ07436"
 repeat_region 37167. 38583
 /note="TIGER1 repeat: matches 2. .1394 of consensus"
 repeat_region 38584. 38937
 /note="THE1B repeat: matches 3. .364 of consensus"
 repeat_region 38938. 40548
 /note="THE1B-INTERNAL repeat: matches 1. .1580 of consensus"
 repeat_region 40551. 40904
 /note="THE1B repeat: matches 3. .364 of consensus"
 repeat_region 40905. 41859
 /note="TIGER1 repeat: matches 1394. .2418 of consensus"
 misc_feature complement(41368. 41893)
 /note="match: GSS: Em:B18147"
 misc_feature complement(41562. 41939)
 /note="match: GSS: Em:AQ074435"
 misc_feature complement(41582. 41936)
 /note="match: GSS: Em:B14391; match: STS: Em:B14391"

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```

repeat_region 26964..27296
/Note="LRR16A repeat: matches 95. .450 of consensus"
27297..27643
/Note="HERVL repeat: matches 3587. .3946 of consensus"
28215..28338
/Note="MSTA repeat: matches 284. .422 of consensus"
28589..29009
/Note="match: GSS: Em:AQ308663"
28617..28812
/Note="MSTA repeat: matches 1. .178 of consensus"
29010..29383
/Note="MLT1B repeat: matches 1. .388 of consensus"
30078..30297
/Note="LRR16A repeat: matches 15. .229 of consensus"
32881..33162
/Note="LIMC5 repeat: matches 7649. .7926 of consensus"
33433..33616
/Note="MIR repeat: matches 40. .260 of consensus"
complement(33617..33834)
/Note="match: GSS: Em:AQ240281"
33899..34036
/Note="MLT11 repeat: matches 269. .404 of consensus"
37901..38084
/Note="AluSg/x repeat: matches 83. .264 of consensus"
38096..38119
/Note="12 copies 2 mer ac 95% conserved"
38855..38952
/Note="MIR repeat: matches 85. .184 of consensus"
39820..40101
/Note="AluY repeat: matches 1. .299 of consensus"
40284..41588
/Note="LTPA7 repeat: matches 2138. .3447 of consensus"
41588..44294
/Note="LTPA7 repeat: matches 3441. .6142 of consensus"
45641..46728
/Note="LIME1 repeat: matches 4994. .6109 of consensus"
46737..47808
/Note="TIGER1 repeat: matches 1350. .2418 of consensus"
47815..48092
/Note="Charlieb repeat: matches 70. .386 of consensus"
48179..48335
/Note="LIPB1 repeat: matches 5999. .6155 of consensus"
48338..48369
/Note="16 copies 2 mer aa 93% conserved"
48370..48882
/Note="LIM4 repeat: matches 4272. .4790 of consensus"
48927..49175
/Note="LIMEC repeat: matches 2146. .2412 of consensus"
49180..49590
/Note="LIMD3 repeat: matches 7310. .7739 of consensus"
49630..49717
/Note="LIMC4 repeat: matches 7219. .7313 of consensus"
49732..50676
/Note="LIMEC repeat: matches 1514. .2152 of consensus"
50716..50836
/Note="LIMEC repeat: matches 1287. .1413 of consensus"
50849..51871
/Note="LIMC1 repeat: matches 5287. .6316 of consensus"
51874..52239
/Note="LIMEC repeat: matches 929. .1302 of consensus"
52268..52519
/Note="LIMEC repeat: matches 523. .780 of consensus"
53098..53206
/Note="Alu repeat: matches 197. .305 of consensus"
53208..53388
/Note="MIR repeat: matches 41. .231 of consensus"
53414..53830
/Note="LTPA8 repeat: matches 5746. .6163 of consensus"
54168..54264
/Note="MIR repeat: matches 2. .111 of consensus"
misc_feature complement(54484..55028)
/Note="match: GSS: Em:AQ318586"
55956..56266
/Note="AluSx repeat: matches 1. .312 of consensus"
56471..56682
/Note="MIR repeat: matches 6. .232 of consensus"
56744..56947
/Note="L2 repeat: matches 2246. .2448 of consensus"
57428..57463
/Note="18 copies 2 mer tt 100% conserved"
complement(57599..58119)
/Note="match: GSS: Em:AQ346372"
58770..59081
/Note="AluY repeat: matches 1. .309 of consensus"
59703..60000
/Note="AluY repeat: matches 1. .294 of consensus"
60141..60194
/Note="L2 repeat: matches 2652. .2706 of consensus"
61142..61188
/Note="L2 repeat: matches 2652. .2698 of consensus"
61806..61910
/Note="L2 repeat: matches 2652. .2698 of consensus"
Query Match 4.3%; Score 26; DB 93; Length 105137;
Best Local Similarity 100.0%; Pred.No. 0.0013;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 85 tgctactgctatcgaatgcataag 110
DB 66659 TGCTACTGCATCTAATGCATAGG 66684

RESULT 4
AC012443 154969 bp DNA HTG 03-APR-2001
LOCUS HOMO sapiens chromosome UNK clone RP11-17G11, WORKING DRAFT
DEFINITION SEQUENCE, 2 unchromed pieces.
AC012443
AC012443.7 GI:13518224
VERSION HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEFIN.
KEYWORDS human.
SOURCE Homo sapiens
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 154969)
Waterston,R.H.
The sequence of Homo sapiens clone
Unpublished
2 (bases 1 to 154969)
Waterston,R.H.
Direct Submission
Submitted (27-OCT-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Apr 3, 2001 this sequence version replaced gi:13446343.

COMMENT
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_NH0017G11
----- Summary Statistics -----
Sequencing vector: M13; 34%
Sequencing vector: plasmid; 53%
Chemistry: Dye-primer ET; 34% of reads
Chemistry: Dye-terminator Big Dye; 53% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 153554 bases at least Q40
Consensus quality: 153754 bases at least Q30
Consensus quality: 153862 bases at least Q20
Insert size: 157000; agarose-ff
Insert size: 153970; sum-of-contigs
Quality coverage: 7.72 in Q20 bases; sum-of-contigs
Quality coverage: 8.02 in Q20 bases; sum-of-contigs
-----
* NOTE: This is a 'working draft' sequence. It currently

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* consists of 2 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

1 153970: contig of 153970 bp in length
 153971 154070: gap of unknown length
 154071 154969: contig of 899 bp in length.

FEATURES

Location/Qualifiers
 1. 154969
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="XUNK"
 /clone="RP11-17G11"

misc_feature

1. 153970
 /note="assembly_name:Contig23
 clone_end:SP6
 vector_side:right"

misc_feature

154071..154969
 /note="assembly_name:Contig5"

BASE COUNT 48393 a 31230 c 30041 g 44839 t 466 others

ORIGIN

Query Match 4.1%; Score 25; DB 62; Length 154969;
 Best Local Similarity 100.0%; Pred. No. 0.0051;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 75 qtgggaacatgctactggcatcta 99

Db 13126 GTGGGACAGCTACTGGCATCTA 13150

RESULT 5

AC090051 183893 bp DNA HTG 04-MAR-2001
 LOCUS Homo sapiens chromosome 12q clone RP11-467E3, WORKING DRAFT
 DEFINITION SEQUENCE, 28 unordered pieces.

ACCESSION

AC090051.2 GI:13194926

KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.

SOURCE

human.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 183893)
 Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-Osman,F.R., Allen,C.,
 Alsbrooks,S.L., Amaralunge,H.C., Are,J.R., Banks,T., Barbara,J.,
 Benton,J., Blimage,K., Blankenburg,K., Bonnin,D., Bouck,J.,
 Bowie,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C.,
 Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C., Carron,T.F.,
 Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R.,
 Chen,Z., Chowdhry,I., Christopoulos,C., Cleveland,C.D., Cox,C.,
 Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C.,
 Denny,A.L., Ding,Y., Dinh,H.H., Douthwaite,K.J., Draper,H.,
 Dugan-Rocha,S., Durbin,K.J., Earnhart,C., Edgar,D., Edwards,C.C.,
 Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J.,
 Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T.,
 Garza,N., Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S.,
 Hamilton,K., Harris,K., Hart,M., Havlak,P., Hawes,A.,
 Hernandez,J., Hernandez,O., Hodgson,A., Hoques,M., Holloway,C.,
 Hollins,B., Homsif,F., Howard,S., Huber,J., Hulyk,S., Hume,J.,
 Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolivet,S.,
 Joudah,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvah,J.,
 Kovar,C., Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C.,
 Lewis,L., Li,J., Li,Z., Lichtarge,O., Lieu,C., Liu,J., Liu,W.,
 Loulseghe,H., Lozano,R.J., Lu,X., Lucier,A., Lucindale,A.,
 Ma,J., Maheshwari,M., Mapua,P., Martin,R., Meador,M.,
 Martinez,E., Massey,E., Mawhiney,E., McLeod,M.P., Meador,M.,
 Mei,G., Metzker,M., Miner,G., Miner,Z., Mitchell,T., Mohabbat,K.,

Morgan,M., Morris,S., Moser,M., Neal,D., Newton,J., Newton,N.,
 Nguyen,A., Nguyen,N., Nickerson,E., Nwokenkwo,S.,
 Oguh,M., Okwuonu,G., Oragunye,N., Oviado,R., Pace,A., Payton,B.,
 Peery,J., Perez,L., Peters,L., Pickens,K., Primus,E., Pu,L.I.,
 Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojibokan,I., Rolfe,M.,
 Ruiz,S., Savary,G., Scherer,S., Scott,G., Shen,H., Shoostari,N.,
 Sisson,I., Sodergren,E., Sonaike,T., Sparks,A., Stanley,K.,
 Stone,H., Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Thomas,N.,
 Tang,H., Tansey,J., Taylor,C., Taylor,T., Telford,B., Vinson,R.,
 Thomas,S., Usmani,K., Vasquez,L., Vera,V., Villalón,D.,
 Walli,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C.,
 Watlington,S., Williams,G., Williamson,A., Wleczyk,R., Wooden,S.,
 Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,
 and Gibbs,R.

Direct Submission
 Unpublished
 2 (bases 1 to 183893)
 Worley,K.C.

Direct Submission

Submitted (12-FEB-2001) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 On Mar 4, 2001 this sequence version replaced gi:12746498.

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: HCVV

Center clone name: RP11-467E3

----- Summary Statistics

Sequencing vector: M13; L08821

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.990329

Consensus quality: 163729 bases at least Q40

Consensus quality: 171892 bases at least Q30

Consensus quality: 175386 bases at least Q20

Estimated insert size: 174626; sum-of-contigs estimation

Quality coverage: 0x in Q20 bases; agarose-gel estimation

Quality coverage: 3.8x in Q20 bases; sum-of-contigs estimation

----- NOTE: Estimated insert size may differ from sequence length

(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 28 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1 25049: contig of 25049 bp in length

* 25149: gap of unknown length

* 43366: contig of 20217 bp in length

* 43667: gap of unknown length

* 45467: contig of 19764 bp in length

* 65231: gap of unknown length

* 75279: contig of 9949 bp in length

* 75379: gap of unknown length

* 85319: contig of 9940 bp in length

* 85320: gap of unknown length

* 95759: contig of 9394 bp in length

* 105252: gap of unknown length

* 105353: contig of 9779 bp in length

* 115231: gap of unknown length

* 123167: contig of 7935 bp in length

* 123266: gap of unknown length

* 130454: contig of 7188 bp in length

* 130554: gap of unknown length

* 134611: contig of 4057 bp in length

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University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Aug 31, 2000 this sequence version replaced gi:9931950.

COMMENT

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_NH0150L12
----- Summary Statistics -----
----- Sequencing vector: M13: 100%
Sequencing vector: plasmid: 0%
Chemistry: Dye-primer; 100% of reads
Chemistry: Dye-terminator; Big Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 184786 bases at least Q40
Consensus quality: 188183 bases at least Q30
Consensus quality: 189812 bases at least Q20
Insert size: 188000; agarose-fp
Insert size: 193094; sum-of-contigs
Quality coverage: 5.74 in Q20 bases; sum-of-contigs
Quality coverage: 5.64 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 13 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 1277: contig of 1277 bp in length
1278 1377: gap of unknown length
1378 11800: contig of 10423 bp in length
1378 11800: contig of 10423 bp in length
11801 11900: gap of unknown length
11901 13553: contig of 1453 bp in length
13554 13453: gap of unknown length
13454 19171: contig of 5718 bp in length
19172 19271: gap of unknown length
19272 32004: contig of 12733 bp in length
32005 32104: gap of unknown length
32105 43844: contig of 11740 bp in length
43845 43945: gap of unknown length
43946 53017: contig of 9073 bp in length
53018 53117: gap of unknown length
53118 64048: contig of 10931 bp in length
64049 64149: gap of unknown length
64150 77841: contig of 13693 bp in length
77842 77941: gap of unknown length
77942 95865: contig of 17924 bp in length
95866 95966: gap of unknown length
95967 122392: contig of 26427 bp in length
122393 122493: gap of unknown length
122494 15346: contig of 31054 bp in length
15347 153646: gap of unknown length
153647 194294: contig of 40648 bp in length.

FEATURES

Source
1. 194294
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="UNK"
/clone="RP11-150L12"
1. 1277
/note="assembly_name:Contig11
clone_end:SP6
vector_side:left"
1378. 11800
/note="assembly_name:Contig15
clone_end:T7
vector_side:right"
11901. 13353
/note="assembly_name:Contig12"

misc_feature

misc_feature

misc_feature

134612 134711: gap of unknown length
134712 138208: contig of 3497 bp in length
138209 138308: gap of unknown length
138309 142509: contig of 4201 bp in length
142510 142609: gap of unknown length
142610 145816: contig of 3207 bp in length
145817 145916: gap of unknown length
145917 145917: contig of 2657 bp in length
145918 148574: gap of unknown length
148575 148673: gap of unknown length
148674 152584: contig of 3911 bp in length
152585 152684: gap of unknown length
152686 156305: contig of 3621 bp in length
156306 156405: gap of unknown length
156406 159712: contig of 3307 bp in length
159713 159812: gap of unknown length
159813 161776: contig of 1964 bp in length
161777 161876: gap of unknown length
161877 165060: contig of 3184 bp in length
165061 165160: gap of unknown length
165161 168260: contig of 3100 bp in length
168261 168361: gap of unknown length
168362 171531: contig of 3171 bp in length
171532 171631: gap of unknown length
171632 174025: contig of 2394 bp in length
174026 174125: gap of unknown length
174126 176898: contig of 2773 bp in length
176899 176998: gap of unknown length
176999 179129: contig of 2131 bp in length
179130 179229: gap of unknown length
179230 181601: contig of 2372 bp in length
181602 181701: gap of unknown length
181702 182782: contig of 1081 bp in length
182783 182882: gap of unknown length
182883 183893: contig of 1011 bp in length.

FEATURES

Source
1. 183893
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="12q"
/clone="RP11-467E3"
54634 a 36554 c 36547 g 52928 t 3230 others

BASE COUNT 54634 a 36554 c 36547 g 52928 t 3230 others

Query Match 4.1%; Score 25; DB 77; Length 183893;
Best Local Similarity 100.0%; Pred. No. 0.0051;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 86 gctactggcatctaatgcatagagg 110
|||||
Db 37859 GCTACTGGCATCTAATGCGATAGAG 37883

RESULT 6
AC079344 194294 bp DNA HTG 31-AUG-2000
LOCUS
DEFINITION Homo sapiens chromosome UNK clone RP11-150L12, WORKING DRAFT
SEQUENCE, 13 chrondomed pieces.

AC079344
VERSION AC079344.2 GI:9954828
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 194294)

Waterston, R.H.

The sequence of Homo sapiens clone

Unpublished

REFERENCE 2 (bases 1 to 194294)

Waterston, R.H.

Direct Submission

TITLE Submitted (28-AUG-2000) Genome Sequencing Center, Washington

JOURNAL


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* 20359 20458: gap of 100 bp
* 20459 21188: contig of 730 bp in length
* 21189 21288: gap of 100 bp
* 21289 22004: contig of 716 bp in length
* 22005 22104: gap of 100 bp
* 22105 22805: contig of 701 bp in length
* 22806 22905: gap of 100 bp
* 22906 23633: contig of 728 bp in length
* 23634 23733: gap of 100 bp
* 23734 24453: contig of 720 bp in length
* 24454 24553: gap of 100 bp
* 24554 25281: contig of 728 bp in length
* 25282 25381: gap of 100 bp
* 25382 26107: contig of 726 bp in length
* 26108 26207: gap of 100 bp
* 26208 26925: contig of 718 bp in length
* 26926 27025: gap of 100 bp
* 27026 27748: contig of 723 bp in length
* 27749 27848: gap of 100 bp
* 27849 28576: contig of 728 bp in length
* 28577 28676: gap of 100 bp
* 28677 29417: contig of 741 bp in length
* 29418 29517: gap of 100 bp
* 29518 30229: contig of 712 bp in length
* 30230 30329: gap of 100 bp
* 30330 31033: contig of 704 bp in length
* 31034 31133: gap of 100 bp
* 31134 31844: contig of 711 bp in length
* 31845 31944: gap of 100 bp
* 31945 32640: contig of 696 bp in length
* 32641 32740: gap of 100 bp
* 32741 33442: contig of 702 bp in length
* 33443 33542: gap of 100 bp
* 33543 34261: contig of 719 bp in length
* 34262 34361: gap of 100 bp
* 34362 35079: contig of 718 bp in length
* 35080 35179: gap of 100 bp
* 35180 35913: contig of 734 bp in length
* 35914 36013: gap of 100 bp
* 36014 36743: contig of 730 bp in length
* 36744 36843: gap of 100 bp
* 36844 37575: contig of 732 bp in length
* 37576 37675: gap of 100 bp
* 37676 38393: contig of 718 bp in length
* 38394 38493: gap of 100 bp
* 38494 39213: contig of 720 bp in length
* 39214 39313: gap of 100 bp
* 39314 40023: contig of 710 bp in length
* 40024 40123: gap of 100 bp
* 40124 40846: contig of 723 bp in length
* 40847 40946: gap of 100 bp
* 40947 41641: contig of 695 bp in length
* 41642 41741: gap of 100 bp
* 41742 42459: contig of 718 bp in length
* 42460 42559: gap of 100 bp
* 42560 43258: contig of 699 bp in length
* 43259 43358: gap of 100 bp
* 43359 44090: contig of 732 bp in length
* 44091 44190: gap of 100 bp
* 44191 44920: contig of 730 bp in length
* 44921 45020: gap of 100 bp
* 45021 45739: contig of 719 bp in length
* 45740 45839: gap of 100 bp
* 45840 46561: contig of 722 bp in length
* 46562 46661: gap of 100 bp
* 46662 47378: contig of 717 bp in length
* 47379 47478: gap of 100 bp
* 47479 48194: contig of 716 bp in length
* 48195 48294: gap of 100 bp
* 48295 49024: contig of 730 bp in length
* 49025 49124: gap of 100 bp
* 49125 49845: contig of 721 bp in length
* 49846 49945: gap of 100 bp

* 49946 50658: contig of 713 bp in length
* 50659 50758: gap of 100 bp
* 50759 51472: contig of 714 bp in length
* 51473 51572: gap of 100 bp
* 51573 52284: contig of 712 bp in length
* 52285 52384: gap of 100 bp
* 52385 53101: contig of 717 bp in length
* 53102 53201: gap of 100 bp
* 53202 53935: contig of 734 bp in length
* 53936 54035: gap of 100 bp
* 54036 54767: contig of 732 bp in length
* 54768 54867: gap of 100 bp
* 54868 55587: contig of 720 bp in length
* 55588 55687: gap of 100 bp
* 55688 56419: contig of 732 bp in length
* 56420 56519: gap of 100 bp

Query Match 3.88; Score 23; DB 78; Length 67084;
Best Local Similarity 100.0%; Pred. No. 0.075;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 163 agaattatctagcccaaatgctc 185
|||||
Db 33966 AGAATTATCTAGCCCCCAATGTC 33944

RESULT 8
LOCUS HS106C24 93959 bp DNA PRI 23-NOV-1999
DEFINITION Human DNA sequence from PAC 106C24, between markers DXS294 and
DSX730 on chromosome X.
ACCESSION 283313
VERSION 283313.1 GI:1730462
KEYWORDS X.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 93959)
Islerwood, J.
Direct Submission
Submitted (02-DEC-1996) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
requests: clonerequest@sanger.ac.uk
IMPORTANT: This sequence is not the entire insert of clone 106C24.
It may be shorter because we only sequence overlapping sections
once, or longer because we arrange for a small overlap between
neighbouring submissions.
This sequence has been finished according to sequence map criteria
as follows. An attempt is made to resolve all sequencing problems,
such as compressions and repeats, but not necessarily within known
annotated human repeat sequence elements (e.g. Alu). Where the
sequence is ambiguous, there is an annotation using the 'unsure'
feature key.
The true left end of clone 106C24 is at 1 in this sequence. The
true right end of clone 20609 is at 47751.
106C24 is from a whole genome PAC library.
location/Qualifiers
source
1. .93959
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="X"
/map="X"
/clone_lib="RPC1-1"
/clone="rp1-106C24"
552. .799
/note="MIR repeat: matches 259. .3 of consensus"
repeat_region
2498. .2582
/note="MIR repeat: matches 57. .145 of consensus"
repeat_region
3438. .3504
/note="MIR repeat: matches 122. .56 of consensus"
repeat_region
4744. .4938
/note="MER3 repeat: matches 2. .200 of consensus"

```

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```

repeat_region 5365..5599
/Note="MIR repeat: matches 8. .236 of consensus"
repeat_region 7810..7942
/Note="MIR repeat: matches 13. .154 of consensus"
repeat_region 8410..8520
/Note="MIR repeat: matches 11. .119 of consensus"
repeat_region 9006..9089
/Note="MIR2 repeat: matches 63. .146 of consensus"
repeat_region 9575..9767
/Note="L1 repeat: matches 5020. .4829 of consensus"
repeat_region 10196..10275
/Note="L1 repeat: matches 106. .4 of consensus"
repeat_region 10515..10619
/Note="MIR repeat: matches 49. .153 of consensus"
repeat_region 11164..11350
/Note="L1 repeat: matches 702. .907 of consensus"
repeat_region 12097..12302
/Note="MIR repeat: matches 30. .262 of consensus"
repeat_region 12338..13023
/Note="MIR2 repeat: matches 146. .64 of consensus"
repeat_region 14787..14844
/Note="29 copies of 2 mer 90 % conserved"
repeat_region 17057..17086
/Note="10 copies of 3 mer 93 % conserved"
repeat_region 17105..17343
/Note="MIR repeat: matches 258. .11 of consensus"
repeat_region 17750..18041
/Note="AluX repeat: matches 297. .5 of consensus"
repeat_region 19193..19360
/Note="MIR repeat: matches 213. .49 of consensus"
repeat_region 20826..20962
/Note="L1 repeat: matches 1. .136 of consensus"
repeat_region 20964..21104
/Note="L1 repeat: matches 92. .234 of consensus"
repeat_region 21093..21869
/Note="L1 repeat: matches 5. .772 of consensus"
repeat_region 21856..26363
/Note="L1 repeat: matches 887. .5390 of consensus"
repeat_region 26214..27104
/Note="L1 repeat: matches 1. .891 of consensus"
repeat_region 29529..29582
/Note="MIR repeat: matches 66. .120 of consensus"
repeat_region 30103..30321
/Note="MIR2 repeat: matches 374. .156 of consensus"
repeat_region 30339..30589
/Note="L1 repeat: matches 909. .659 of consensus"
repeat_region 30596..30743
/Note="MIR repeat: matches 149. .1 of consensus"
repeat_region 30834..31103
/Note="MIR repeat: matches 1231. .961 of consensus"
repeat_region 31580..31951
/Note="MIR repeat: matches 599. .231 of consensus"
repeat_region 31957..32266
/Note="AluX repeat: matches 297. .1 of consensus"
repeat_region 32267..32482
/Note="MIR repeat: matches 249. .5 of consensus"
repeat_region 32487..32545
/Note="MIR2 repeat: matches 374. .313 of consensus"
repeat_region 32551..32850
/Note="L1 repeat: matches 1. .301 of consensus"
repeat_region 32870..32911
/Note="MIR repeat: matches 270. .312 of consensus"
repeat_region 32912..33202
/Note="MIR repeat: matches 297. .1 of consensus"
repeat_region 33703..33777
/Note="MIR repeat: matches 345. .271 of consensus"
repeat_region 33811..33871
/Note="MIR repeat: matches 990. .1050 of consensus"
repeat_region 33898..34115
/Note="MIR repeat: matches 210. .1 of consensus"
repeat_region 35085..35213
/Note="MIR repeat: matches 152. .29 of consensus"
35255..35321
/Note="MIR repeat: matches 149. .84 of consensus"
35621..36097
/Note="MIR repeat: matches 1. .465 of consensus"
36588..36721
/Note="MIR repeat: matches 119. .256 of consensus"
38850..39742
/Note="L1 repeat: matches 893. .1 of consensus"
39598..44765
/Note="L1 repeat: matches 5390. .386 of consensus"
45104..45334
/Note="L1 repeat: matches 255. .12 of consensus"
48462..48649
/Note="L1 repeat: matches 935. .737 of consensus"
50125..53643
/Note="L1 repeat: matches 1877. .5390 of consensus"
53494..54383
/Note="L1 repeat: matches 1. .891 of consensus"
54743..55195
/Note="L1 repeat: matches 867. .393 of consensus"
55530..56049
/Note="L1 repeat: matches 889. .311 of consensus"
57142..57399
/Note="L1 repeat: matches 5129. .5389 of consensus"
57251..58278
/Note="L1 repeat: matches 1. .1055 of consensus"
58483..58557
/Note="L1 repeat: matches 815. .889 of consensus"
58564..58597
/Note="L1 repeat: matches 2 mer 82 % conserved"
58905..59344
/Note="L1 repeat: matches 12. .484 of consensus"
59359..59409
/Note="L1 repeat: matches 1488. .1538 of consensus"
59440..59664
/Note="MIR repeat: matches 230. .1 of consensus"
60853..60918
/Note="MIR repeat: matches 78. .143 of consensus"
61282..61335
/Note="MIR repeat: matches 2 mer 96 % conserved"
61374..61454
/Note="MIR repeat: matches 146. .64 of consensus"
61902..62103
/Note="MIR repeat: matches 9. .247 of consensus"
64007..64146
/Note="MIR repeat: matches 32. .170 of consensus"
64287..64532
/Note="MIR repeat: matches 70. .324 of consensus"
67972..68217
/Note="MIR repeat: matches 4. .247 of consensus"
68347..68471
/Note="MIR repeat: matches 15. .144 of consensus"
70707..70748
/Note="MIR repeat: matches 88. .47 of consensus"
71075..71375
/Note="AluX repeat: matches 1. .300 of consensus"
71608..71649
/Note="MIR repeat: matches 141. .100 of consensus"
71736..71944
/Note="MIR repeat: matches 5. .215 of consensus"
71985..72402
/Note="MIR repeat: matches 2. .426 of consensus"
72407..72502
/Note="MIR repeat: matches 233. .328 of consensus"
72503..72804
/Note="MIR repeat: matches 301. .2 of consensus"
72806..72876
/Note="MIR repeat: matches 324. .394 of consensus"
72877..72979
/Note="MIR repeat: matches 1. .102 of consensus"
72909..73456
/Note="MIR repeat: matches 32. .593 of consensus"

```

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* of the gaps between them are based on estimates that have
 * provided by the submitter.
 * This sequence will be replaced
 * by the finished sequence as soon as it is available and
 * the accession number will be preserved.
 * the accession number will be preserved.
 * 1 103757: contig of 103757 bp in length.

FEATURES

source
 1. 103757
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="15"
 /map="15q15"
 /clone="CTD-2190J1"
 /clone_lib="Cal Tech human BAC library D"
 /note="This clone overlaps RP11-213B23 and RP11-616K22"
 /note="This clone overlaps RP11-213B23 and RP11-616K22"

BASE COUNT 32585 a 21521 c 21252 g 28399 t
 ORIGIN

Query Match 3.8%; Score 23; DB 75; Length 103757;
 Best Local Similarity 100.0%; Pred. No. 0.077;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 163 agaattatctagcccaaatgctc 185
 |||||

Db 94438 AGAATTATCTAGCCCAATGTC 94416

RESULT 10

AC002554 127590 bp DNA PRI 27-AUG-1998
 LOCUS Human Chromosome 1p11.2 PAC clone pDJ404m15, complete sequence.

AC002554
 DEFINITION

AC002554
 ACCESSION

AC002554.1 GI:3478647

VERSION

HTG.

KEYWORDS

human.

SOURCE

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 127590)

Evans, G.A., Athanasiou, M., Aguayo, P., Armstrong, D., Basit, M.,

Buettner, J., Bumeister, R., Card, P., deSailboat, F., Dunn, J.,

English, C., Ethridge, S., Garner, H.R., Gee, V., Gordon, M., Gotway, G.,

Grant, O., Hahner, L., Harris, J., Hinson, S.,

Major, T., McFarland, J., Newton, J., Osborne-Lawrence, S.,

Schageman, J., Schultz, R.A., Stimson, S., Syed, M. and Ward, T.

HTGS Submission

Unpublished

2 (bases 1 to 127590)

Evans, G.A., Athanasiou, M., Bradbury, P., Brignac, S., Bumeister, R.,

Davis, C., English, C., Franklin, T.L., Garner, H.R., Gordon, M.,

Gotway, G., Grant, O., Hahner, L., Harris, J., Hinson, S.,

Narayanaswamy, U., Newton, J., O'Brien, K., Oliver, T., Patel, P.,

Probst, S., Rayner, S., Schageman, J., Schilling, P., Schultz, R.,

Syed, M., Valenzuela, D., Ward, T. and Wilson, R.

Direct Submission

Submitted (23-SEP-1997) Genome Science and Technology Center,

University of Texas Southwestern Medical Center at Dallas, 5323

Harry Hines Blvd, Dallas, TX 75235-8591, USA

3 (bases 1 to 127590)

Evans, G.A., Athanasiou, M., Basit, M., Bradbury, P., Brignac, S.,

Bumeister, R., Davis, C., English, C., Franklin, T.L., Garner, H.R.,

Gee, V., Gordon, M., Gotway, G., Grant, O., Hahner, L., Harris, J.,

Hinson, S., Narayanaswamy, U., Newton, J., O'Brien, K., Patel, P.,

Schageman, J., Schilling, P., Schultz, R., Syed, M., Valenzuela, D.,

Ward, T. and Wilson, R.

Direct Submission

Submitted (17-DEC-1997) Genome Science & Technology Center,

University of Texas Southwestern Medical Center, 5323 Harry Hines

Bldv, Dallas, TX 75235-8591, USA

4 (bases 1 to 127590)

Evans, G.A., Athanasiou, M., Aguayo, P., Armstrong, D., Basit, M.,

Buettner, J., Bumeister, R., Card, P., deSailboat, F., Dunn, J.,

repeat_region 73468..73517
 /note="25 copies of 2 mer 96 & conserved"
 repeat_region 73543..74432
 /note="MST-INTERNAL repeat: matches 532..1458 of
 consensus"
 repeat_region 74466..75346
 /note="11PAL2 repeat: matches 912..4 of consensus"
 repeat_region 75199..75954
 /note="L1 repeat: matches 5390..767 of consensus"

Query Match 3.8%; Score 23; DB 92; Length 93959;
 Best Local Similarity 100.0%; Pred. No. 0.076;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 163 agaattatctagcccaaatgctc 185
 |||||

Db 85580 AGAATTATCTAGCCCAATGTC 85602

RESULT 9

AC073940/c 103757 bp DNA HTG 14-APR-2001
 LOCUS Homo sapiens chromosome 15 clone CTD-2190J1 map 15q15, ***

AC073940
 DEFINITION

AC073940
 ACCESSION

AC073940.3 GI:13624386

VERSION

HTG; HTGS_PHASE2; HTGS_FULLTOP.

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 103757)

Rowen, L., Madan, A., Qin, S., Baradarani, L., Birditt, B., Bloom, S.,

Burke, J., Dors, M., Fleetwood, P., Kaur, A., Madan, A., Nesbitt, R.,

Pate, D. and Hood, L.

Sequencing of human chromosome 15 D15S146-D15S117 region

Unpublished

2 (bases 1 to 103757)

Rowen, L., Madan, A., Qin, S., Abbasi, N., Baradarani, L., Birditt, B.,

Bloom, S., Dors, M., Dickhoff, R., Fleetwood, P., Harrison, G., Kaur, A.,

Madan, A., Nesbitt, R., Shaffer, T. and Hood, L.

Direct Submission

Submitted (07-JUL-2000) Multimegabase Sequencing Center, Institute

for Systems Biology, 4225 Roosevelt Way NE, Suite 200, Seattle, WA

98105, USA

On Apr 14, 2001 this sequence version replaced gi:12658005.

Center: Genome Center

Center code: UMWSC

Web site: http://chroma.mbt.washington.edu/msg_www

Contact: leetowen@systemsbiology.org

Summary Statistics

Sequencing vector: pUC18; L08752

Chemistry: Dye-terminator Big Dye; 90% of reads

Chemistry: Dye-terminator Big Dye; 10% of reads

Assembly program: Phrap; version 0.990399

Insert size: 100000; agarose-ff

Quality coverage: 10.2x in Q20 bases; sum-of-contigs

Sequence Quality Assessment:

This entry has been annotated with sequence quality

estimates computed by the Phrap assembly program.

All manually edited bases have been reduced to quality zero.

Quality levels above 40 are expected to have less than

1 error in 10,000 bp.

Base-by-base quality values are not generally visible from the

Genbank flat file format but are available as part

of this entry's ASN.1 file.

* NOTE: This is a 'working draft' sequence. It currently

* consists of 1 contigs. Gaps between the contigs

* are represented as runs of N. The order of the pieces

* is believed to be correct as given, however the sizes

TITLE

English,C., Ethridge,S., Garner,H.R., Gee,V., Gordon,M., Gotway,G., Grant,O., Hahner,L., Joslin,J., Lewis,E., Loo,H., Loo,K.N., Major,T., McFarland,J., Newton,J., Osborne-Lawrence,S., Schageman,J., Schultz,R.A., Stimson,S., Syed,M. and Ward,T.

JOURNAL

Direct Submission
Submitted (27-AUG-1998) Genome Science & Technology Center,
University of Texas Southwestern Medical Center, 5323 Harry Hines Blvd., Dallas, TX 75235-8591, USA

COMMENT

On Aug 27, 1998 this sequence version replaced gi:2695561.
CHROMOSOMAL LOCUS: This PAC clone comes from the llpl1.2 region mapped between STS markers D1L578 and D1L580sh.
MARCKER CONFIRMATION: STS sequence confirmed: D1L578
MAPPED CLONE OVERLAP: pdJ404cl0.
IMPORTANT: This submission contains the entire insert of clone pdJ404m15. pdJ404m15 comes from the RPLI-3 PAC library constructed at the Roswell Park Cancer Institute by the Pieter de Jong group. This clone has been finished according to strict quality criteria and attempts have been made to resolve all base calling problems such as compressions and repetitive elements. The expected Phred/Phrap calculated errors/10kb is 0.23. In addition, this sequence has been finished such that 99.9% of consensus base calls consist of either double-stranded coverage or 2 types of labelling chemistry on one strand. Additional information regarding this clone may be found on the GENECC web page at http://genec.swned.edu/llpl1.htm.

FEATURES	Source	Location/Qualifiers
repeat_region	/rpt_family="Alu"	.127590
repeat_region	/organism="Homo sapiens"	.
repeat_region	/db_xref="taxon:9606"	.
repeat_region	complement(1..69)	.
repeat_region	/rpt_family="MER41"	96..385
repeat_region	/rpt_family="Alu"	.
repeat_region	complement(400..821)	.
repeat_region	/rpt_family="MER41"	.
repeat_region	complement(1101..1382)	.
repeat_region	/rpt_family="LTR9"	.
repeat_region	complement(1436..1731)	.
repeat_region	/rpt_family="Alu"	.
repeat_region	complement(1797..1937)	.
repeat_region	/rpt_family="LTR9"	.
repeat_region	complement(3274..3558)	.
repeat_region	/rpt_family="Alu"	.
repeat_region	complement(3923..4100)	.
repeat_region	/rpt_family="MER41"	4779..5469
repeat_region	/rpt_family="LTR8"	.
repeat_region	complement(7262..7513)	.
repeat_region	/rpt_family="Alu"	.
repeat_region	complement(7651..7964)	.
repeat_region	/rpt_family="Alu"	11021..11306
repeat_region	/rpt_family="Alu"	11424..12625
repeat_region	/rpt_family="LI"	13008..13297
repeat_region	/rpt_family="Alu"	13356..13506
repeat_region	/rpt_family="LI"	complement(14032..14334)
repeat_region	/rpt_family="Alu"	14861..15068
repeat_region	/rpt_family="MER20"	15934..16238
repeat_region	/rpt_family="Alu"	16914..17018
repeat_region	/rpt_family="Alu"	complement(17093..17337)
repeat_region	/rpt_family="Alu"	complement(17362..17691)
repeat_region	/rpt_family="MER42"	17918..18156
repeat_region	/rpt_family="Alu"	.

complement(18206..18402)	complement(18206..18402)
rpt_family="MIR"	19740..19807
rpt_family="Alu"	complement(21285..21395)
rpt_family="MLT1"	21587..21762
rpt_family="MLT1"	21779..21846
rpt_family="MLT1"	21857..21924
rpt_family="WSTAR"	23972..24546
rpt_family="MLT1"	24638..24733
rpt_family="MIR"	complement(24860..25148)
rpt_family="Alu"	26031..26203
rpt_family="Alu"	26528..26717
rpt_family="Alu"	complement(27650..27708)
rpt_family="MIR"	complement(29082..29343)
rpt_family="Alu"	30684..30980
rpt_family="Alu"	32634..32818
rpt_family="Alu"	complement(35666..35779)
rpt_family="Alu"	36025..36096
rpt_family="MIR"	36923..37030
rpt_family="MIR"	complement(37101..37189)
rpt_family="THE1"	37653..37940
rpt_family="Alu"	38374..38647
rpt_family="Alu"	complement(38907..39198)
rpt_family="Alu"	complement(40840..40945)
rpt_family="MERS"	complement(41499..41785)
rpt_family="Alu"	42735..43073
rpt_family="MLT1"	complement(43666..43951)
rpt_family="Alu"	44899..45060
rpt_family="MIR"	complement(45617..45784)
rpt_family="Alu"	45878..46030
rpt_family="MIR"	complement(46105..46157)
rpt_family="MLT1"	complement(46177..46305)
rpt_family="MLT1"	46676..46841
rpt_family="MER2"	46871..46959
rpt_family="Alu"	complement(46983..47265)
rpt_family="Alu"	47306..47395
rpt_family="Alu"	47503..47580
rpt_family="Alu"	47627..47741

us-09-656-668-198.oli.rge

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```

repeat_region      /rpt_family="MER2"
complement(47741..47839)
/rpt_family="MIR"
49558..49671
/rpt_family="MIR"
50232..50508
/rpt_family="Alu"
complement(52633..52764)
/rpt_family="MIR"
complement(53356..53634)
/rpt_family="Alu"

Query Match      3.98; Score 23; DB 85; Length 127590;
Best Local Similarity 100.0%; Pred. No. 0.078;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 163 agaattatctagcccaaatctc 185
|||||
Db 15032 AGAATTATCTAGCCCAAAATCTC 15054

RESULT 11
AC067916 134394 bp DNA HTG 09-JUN-2000
LOCUS Homo sapiens chromosome X clone RP11-513J24 map X, *** SEQUENCING
DEFINITION IN PROGRESS ***, 32 unordered pieces.
ACCESSION AC067916
VERSION AC067916.2 GI:8389561
KEYWORDS HTG; HTGS_PHASE1.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 134394)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F.,
Boguslavsky,L., Bouckgalter,B., Brown,A., Burgett,G.,
Campiano,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,
Collamore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,
Dodgson,S., Domino,M., Doyle,M., Ferreira,P., FitzHugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
Grand-pierre,N., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,
Klein,J., LaRoque,K., Lamazares,R., Landers,T., Lehocsky,J.,
Levine,R., Lieu,C., Liu,G., Locke,K., Macdonald,P., Marquis,N.,
McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R.,
Meldrum,J., Meneus,D., Mihova,T., Miranda,C., Mienda,V., Morrow,J.,
Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neill,D., Olivaz,T.M., Oliver,J., Peterson,K., Pierre,N.,
Pisani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
Stange-Roman,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tessaye,S., Theodore,J., Tirrell,A., Travers,M., Trigilio,J.,
Vassiliev,H., Viel,R., Vo.A., Wilson,B., Wu,X., Wyman,D., Ye.W.J.,
Young,G., Zainoun,J., Zimmer,A. and Zody,M.

Direct Submission
Submitted (27-APR-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jun 9, 2000 this sequence version replaced gi:7652204.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L6355

```

Center clone name: 513_J_24

NOTE: This is a 'working draft' sequence. It currently consists of 32 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1240: contig of 1240 bp in length
1241 1340: gap of 100 bp
1341 2613: contig of 1273 bp in length
2614 2713: gap of 100 bp
2714 4159: contig of 1446 bp in length
4160 4259: gap of 100 bp
4260 5435: contig of 1176 bp in length
5436 5535: gap of 100 bp
5536 6901: contig of 1366 bp in length
6902 7001: gap of 100 bp
7002 8119: contig of 1118 bp in length
8120 8219: gap of 100 bp
8220 10213: contig of 1994 bp in length
10214 10313: gap of 100 bp
10314 12093: contig of 1780 bp in length
12094 12193: gap of 100 bp
12194 13509: contig of 1316 bp in length
13510 13609: gap of 100 bp
13610 14871: contig of 1262 bp in length
14872 14971: gap of 100 bp
14972 16966: contig of 1995 bp in length
16967 17066: gap of 100 bp
17067 18732: contig of 1666 bp in length
18733 18832: gap of 100 bp
18833 20655: contig of 1823 bp in length
20656 20755: gap of 100 bp
20756 22721: contig of 1966 bp in length
22722 22821: gap of 100 bp
22822 24991: contig of 2170 bp in length
24992 25091: gap of 100 bp
25092 25443: contig of 352 bp in length
25444 25543: gap of 100 bp
25544 28606: contig of 3063 bp in length
28607 28706: gap of 100 bp
28707 30875: contig of 2169 bp in length
30876 30975: gap of 100 bp
30976 34223: contig of 3248 bp in length
34224 34323: gap of 100 bp
34324 37911: contig of 3588 bp in length
37912 38011: gap of 100 bp
38012 42033: contig of 4022 bp in length
42034 42133: gap of 100 bp
42134 45055: contig of 2922 bp in length
45056 45155: gap of 100 bp
45156 50793: contig of 5638 bp in length
50794 50893: gap of 100 bp
50894 56638: contig of 5745 bp in length
56639 56738: gap of 100 bp
56739 62187: contig of 5449 bp in length
62188 62288: gap of 100 bp
62289 69769: contig of 7482 bp in length
69770 69869: gap of 100 bp
69870 77546: contig of 7677 bp in length
77547 77646: gap of 100 bp
77647 84823: contig of 7077 bp in length
84824 92916: contig of 8093 bp in length
92917 93016: gap of 100 bp
93017 107613: contig of 14597 bp in length
107614 107713: gap of 100 bp
107714 119925: contig of 12212 bp in length
119926 120025: gap of 100 bp
120026 134394: contig of 14369 bp in length.

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FEATURES
source
Location/Qualifiers
1..134394
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="X"
/map="X"
/clone="RP11-513J24"
/clone.lib="RP11-11 Human Male BAC"
1..1240
/note="assembly_fragment"
1341..2613
/note="assembly_fragment"
2714..4159
/note="assembly_fragment"
4260..5435
/note="assembly_fragment"
5536..6901
/note="assembly_fragment"
7002..8119
/note="assembly_fragment"
8220..10213
/note="assembly_fragment"
10314..12093
/note="assembly_fragment"
12194..13509
/note="assembly_fragment"
13610..14871
/note="assembly_fragment"
14972..16966
/note="assembly_fragment"
17067..18732
/note="assembly_fragment"
18833..20655
/note="assembly_fragment"
20756..22721
/note="assembly_fragment"
22822..24991
/note="assembly_fragment"
25092..25443
/note="assembly_fragment"
clone_end:SP6
vector_side:right"
25544..28606
/note="assembly_fragment"
28707..30875
/note="assembly_fragment"
30976..34223
/note="assembly_fragment"
34324..37911
/note="assembly_fragment"
38012..42033
/note="assembly_fragment"
42134..44505
/note="assembly_fragment"
clone_end:T7
vector_side:right"
45156..50793
/note="assembly_fragment"
50894..56638
/note="assembly_fragment"
56739..62187
/note="assembly_fragment"
62288..69769
/note="assembly_fragment"
69870..77546
/note="assembly_fragment"
77647..84723
/note="assembly_fragment"
84824..92916
/note="assembly_fragment"
93017..107613
/note="assembly_fragment"
107714..119925
/note="assembly_fragment"

misc_feature
120026..134394
/note="assembly_fragment"
misc_feature
120026..134394
/note="assembly_fragment"
BASE COUNT
39328 a 25781 c 26014 g 40164 t 3107 others
Query Match
3.8%; Score 23; DB 73; Length 134394;
Best Local Similarity
100.0%; Pred. No. 0.078;
Matches
23; Conservative
0; Mismatches
0; Indels
0; Gaps
0;

QY 163 agaatattctagcccaaatgtc 185
|||||
Db 77870 AGAATTATCTAGCCCAATGTC 77848

RESULT 12
AC011573 151357 bp DNA HTG 12-MAR-2000
LOCUS Homo sapiens clone RP11-12G12, WORKING DRAFT SEQUENCE, 28 unordered
DEFINITION pieces.
AC011573
AC011573.3 GI:7230070
VERSION HTG; HTGS_PHASE1; HTGS_DRAFT.
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 151357)
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
Homo sapiens, clone RP11-12G12
Unpublished
2 (bases 1 to 151357)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Boukhgalter,B.,
Brown,A., Castelle,A., Colangelo,M., Collins,S., Collymore,A.,
Cooke,P., Deatellano,K., Dewar,K., Domino,M., Donegan,L., Doyle,M.,
Ferrelira,P., Fitzhugh,W., Forrest,C., Funke,R., Gage,D., Horton,L.,
Galagan,J., Gardyna,S., Grant,G., Hagos,B., Hearford,A., Klein,J.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Marquis,N.,
Lehoczky,J., Lieu,C., Locke,K., McKernan,K., McLaughlin,J., Meldrim,J.,
McEwan,P., McGurk,A., McKernan,C.H., O'Connor,T., O'Donnell,P.,
Morrow,J., Naylor,J., Norman,C.H., Roy,A., Santos,R., Severy,P.,
Peterson,K., Pollara,V., Riley,R., Riley,C., Subramanian,A., Talamas,J.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tefaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.
Direct Submission
Submitted (07-OCT-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Mar 12, 2000 this sequence version replaced gi:6249718.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www.seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
----- Project Information
Center project name: L3265
Center clone name: 12.G.12
----- Summary Statistics
Sequencing vector: M13; M7815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 101368 bases at least Q40
Consensus quality: 127052 bases at least Q30
Consensus quality: 139101 bases at least Q20
Insert size: 147000; agarose-fp
Insert size: 148657; sum-of-contigs
Quality coverage: 3.5 in Q20 bases; agarose-fp
Quality coverage: 3.5 in Q20 bases; sum-of-contigs
-----
* NOTE: This is a 'working draft' sequence. It currently

```

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* consists of 28 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 1505: contig of 1505 bp in length
* 1506 1605: gap of 100 bp
* 1606 2995: contig of 1390 bp in length
* 2996 3095: gap of 100 bp
* 3096 4668: contig of 1573 bp in length
* 4669 4768: gap of 100 bp
* 4769 6111: contig of 1343 bp in length
* 6112 6211: gap of 100 bp
* 6212 7559: contig of 1348 bp in length
* 7560 7659: gap of 100 bp
* 7660 8904: contig of 1245 bp in length
* 8905 9004: gap of 100 bp
* 9005 10441: contig of 1437 bp in length
* 10442 10541: gap of 100 bp
* 10542 12992: contig of 2451 bp in length
* 12993 13092: gap of 100 bp
* 13093 16001: contig of 2909 bp in length
* 16002 16101: gap of 100 bp
* 16102 18808: contig of 2707 bp in length
* 18809 18908: gap of 100 bp
* 18909 22389: contig of 3481 bp in length
* 22390 22489: gap of 100 bp
* 22490 25729: contig of 3240 bp in length
* 25730 25829: gap of 100 bp
* 25830 28948: contig of 3119 bp in length
* 28949 29048: gap of 100 bp
* 29049 32582: contig of 3534 bp in length
* 32583 32682: gap of 100 bp
* 32683 36104: contig of 3422 bp in length
* 36105 36204: gap of 100 bp
* 36205 39825: contig of 3621 bp in length
* 39826 39925: gap of 100 bp
* 39926 45184: contig of 5259 bp in length
* 45185 45284: gap of 100 bp
* 45285 50916: contig of 5632 bp in length
* 50917 51016: gap of 100 bp
* 51017 56385: contig of 5369 bp in length
* 56386 56485: gap of 100 bp
* 56486 63439: contig of 6954 bp in length
* 63440 63539: gap of 100 bp
* 63540 70246: contig of 6707 bp in length
* 70247 70346: gap of 100 bp
* 70347 76319: contig of 5973 bp in length
* 76320 76419: gap of 100 bp
* 76420 85131: contig of 8712 bp in length
* 85132 85231: gap of 100 bp
* 85232 91974: contig of 6743 bp in length
* 91975 92074: gap of 100 bp
* 92075 102742: contig of 10668 bp in length
* 102743 102842: gap of 100 bp
* 102843 110539: contig of 7697 bp in length
* 110540 110639: gap of 100 bp
* 110640 119486: contig of 8847 bp in length
* 119487 119586: gap of 100 bp
* 119587 151357: contig of 31771 bp in length.

FEATURES
source

1. .151357
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/db_xref="taxon:9606"
/clone="RP11-12G12"
/clone_lib="RPC1-11 Human Male BAC"
1. .1505
/note="assembly_fragment"
1606. .2995
/note="assembly_fragment"
3096 4668
misc_feature
misc_feature
misc_feature

misc_feature /note="assembly_fragment"
4769. .6111
/note="assembly_fragment"
6212. .7559
/note="assembly_fragment"
7660. .8904
/note="assembly_fragment"
9005. .10441
/note="assembly_fragment"
10542. .12992
/note="assembly_fragment"
13093. .16001
/note="assembly_fragment"
16102. .18808
/note="assembly_fragment"
18909. .22389
/note="assembly_fragment"
22490. .25729
/note="assembly_fragment"
25830. .28948
/note="assembly_fragment"
29049. .32582
/note="assembly_fragment"
32683. .36104
/note="assembly_fragment"
36205. .39825
/note="assembly_fragment"
39926. .45184
/note="assembly_fragment"
45285. .50916
/note="assembly_fragment"
51017. .56385
/note="assembly_fragment"
56486. .63439
/note="assembly_fragment"
clone_end:SP6
vector_side:right"
63540. .70246
/note="assembly_fragment"
70347. .76319
/note="assembly_fragment"
76420. .85131
/note="assembly_fragment"
85232. .91974
/note="assembly_fragment"
clone_end:T7
vector_side:left"
92075. .102742
/note="assembly_fragment"
102843. .110539
/note="assembly_fragment"
110640. .119486
/note="assembly_fragment"
119587. .151357
/note="assembly_fragment"
BASE COUNT 40770 a 33565 c 33412 g 40768 t 2842 others
ORIGIN
Query Match 3.8%; Score 23; DB 62; Length 151357;
Best Local Similarity 100.0%; Pred. No. 0.078;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 56 ttttgattgtcacacctgggtgg 78
|||||
Db 56633 TTTTGATTGTACACCTGGTGG 56655
RESULT 13
AC034161 157402 bp DNA HTG 01-SEP-2000
LOCUS Homo sapiens chromosome 11 clone RP11-397J5, WORKING DRAFT
DEFINITION SEQUENCE, 18 unordered pieces.

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AC034161
AC034161.3 GI:9958241
HTGS_PHASE1; HTGS_DRAFT.
human.
Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 157402)
Waterston,R.H.
The sequence of Homo sapiens clone
Unpublished
2 (bases 1 to 157402)
Waterston,R.H.
Direct Submission
Submitted (04-APR-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Sep 1, 2000 this sequence version replaced gi:7523965.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H.NH0397J05
----- Summary Statistics -----
Sequencing vector: M13; 100%
Chemistry: Dye-terminator Big Dye; 0% of reads
Chemistry: Dye-terminator Big Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 149716 bases at least Q40
Consensus quality: 152398 bases at least Q30
Consensus quality: 153456 bases at least Q20
Insert size: 191000; agarose-fp
Insert size: 155702; sum-of-contigs
Quality coverage: 3.43 in Q20 bases; agarose-fp
Quality coverage: 4.25 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 5077: contig of 5077 bp in length
* 5078 5177: gap of unknown length
* 5178 11207: contig of 6030 bp in length
* 11208 11307: gap of unknown length
* 11308 18089: contig of 6782 bp in length
* 18090 18189: gap of unknown length
* 18190 24820: contig of 6631 bp in length
* 24821 24820: gap of unknown length
* 32091 32191: contig of 7171 bp in length
* 32192 45992: gap of unknown length
* 45993 46092: contig of 13801 bp in length
* 46093 52957: gap of unknown length
* 52958 53057: contig of 8865 bp in length
* 53058 61361: gap of unknown length
* 61361 69613: contig of 8153 bp in length
* 69614 69713: gap of unknown length
* 69714 80408: contig of 10695 bp in length
* 80409 80508: gap of unknown length
* 80509 91848: contig of 11340 bp in length
* 91849 91948: gap of unknown length
* 91949 115447: contig of 23499 bp in length
* 115448 115547: gap of unknown length
* 115548 142318: contig of 26771 bp in length
* 142319 142418: gap of unknown length

145665: contig of 3247 bp in length
145765: gap of unknown length
148715: contig of 2950 bp in length
148815: gap of unknown length
151082: contig of 2267 bp in length
151182: gap of unknown length
153839: contig of 2657 bp in length
153939: gap of unknown length
157402: contig of 3463 bp in length.
Location/Qualifiers
1. 157402
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="11"
/clone="RP11-397J5"
1. 5077
/note="assembly_name:Contig10"
5178. 11207
/note="assembly_name:Contig11"
11308. 18089
/note="assembly_name:Contig12"
18190. 24820
/note="assembly_name:Contig13"
24921. 32091
/note="assembly_name:Contig14"
clone_end:SP6
vector_side:right
32192. 45992
/note="assembly_name:Contig19"
clone_end:T7
vector_side:right
46093. 52957
/note="assembly_name:Contig15"
53058. 61360
/note="assembly_name:Contig16"
61461. 69613
/note="assembly_name:Contig17"
69714. 80408
/note="assembly_name:Contig18"
80509. 91848
/note="assembly_name:Contig20"
91949. 115447
/note="assembly_name:Contig21"
115548. 142318
/note="assembly_name:Contig22"
142419. 145665
/note="assembly_name:Contig5"
145766. 148715
/note="assembly_name:Contig6"
148816. 151082
/note="assembly_name:Contig7"
151183. 153839
/note="assembly_name:Contig8"
153940. 157402
/note="assembly_name:Contig9"
BASE COUNT 49815 a 29500 c 29459 g 46922 t 1706 others
ORIGIN

Query Match 3.8%; Score 23; DB 71; Length 157402;
Best Local Similarity 100.0%; Pred. No. 0.078;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 163 aqaattatctagcccaaatgtc 185
|||||
Db 13008 AGAATTACTAGCCCAATGTC 13030
RESULT 14
AC021862 159723 bp DNA HTG 07-JUL-2000
LOCUS Homo sapiens chromosome 11 clone RP11-674E24, WORKING DRAFT
DEFINITION SEQUENCE, 18 unordered pieces.

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AC021862
VERSION
KEYWORDS
SOURCE

AC021862.4 GI:8570373
HTG; HTGS_PHASE1; HTGS_DRAFT.

ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 159723)

Waterston, R.H.

The sequence of Homo sapiens clone

Unpublished

2 (bases 1 to 159723)

Waterston, R.H.

Direct Submission
Submitted (20-JAN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Jun 17, 2000 this sequence version replaced gi:7235315.

COMMENT

----- Genome Center -----

Center: Washington University Genome Sequencing Center

Center code: WUGSC

Web site: http://genome.wustl.edu/gsc/index.shtml

----- Project Information -----

Center project name: H_NH0674E24

----- Summary Statistics -----

Sequencing vector: M13; 76%

Sequencing vector: plasmid; 24%

Chemistry: Dye-primer ET; 76% of reads

Chemistry: Dye-terminator Big Dye; 24% of reads

Assembly program: Phrap; version 0.990319

Consensus quality: 146571 bases at least Q40

Consensus quality: 150482 bases at least Q30

Consensus quality: 152816 bases at least Q20

Insert size: 167000; agarose-fp

Insert size: 158023; sum-of-contigs

Quality coverage: 3.44 in Q20 bases; agarose-fp

Quality coverage: 3.71 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 1195: contig of 1195 bp in length
* 1196 1295: gap of unknown length
* 1296 3879: contig of 2584 bp in length
* 3880 3979: gap of unknown length
* 3980 6597: contig of 2618 bp in length
* 6598 6697: gap of unknown length
* 6698 10040: contig of 3343 bp in length
* 10041 10140: gap of unknown length
* 10141 12973: contig of 2833 bp in length
* 12974 13074: gap of unknown length
* 13074 15987: contig of 2914 bp in length
* 15988 16087: gap of unknown length
* 16088 19014: contig of 2927 bp in length
* 19015 19114: gap of unknown length
* 19115 22164: contig of 3050 bp in length
* 22165 22264: gap of unknown length
* 22265 26798: contig of 4534 bp in length
* 26799 26898: gap of unknown length
* 26899 34198: contig of 7300 bp in length
* 34199 34298: gap of unknown length
* 34299 42840: contig of 8542 bp in length
* 42841 42941: gap of unknown length
* 42941 48833: contig of 5893 bp in length
* 48834 59689: gap of unknown length
* 59690 59789: contig of 10756 bp in length
* 59789 gap of unknown length

59790 72021: contig of 12232 bp in length
* 72022 72121: gap of unknown length
* 72122 87123: contig of 15002 bp in length
* 87124 87223: gap of unknown length
* 87224 106312: contig of 19089 bp in length
* 106313 106412: gap of unknown length
* 106413 133985: contig of 27573 bp in length
* 133986 134086: gap of unknown length
* 134086 159723: contig of 25638 bp in length.
FEATURES
source
Location/Qualifiers
1..159723
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/db_xref="taxon:9606"
/chromosome="11"
/clone="RP11-674E24"
1..1195
misc_feature
/note="assembly_name:Contig7"
1296..3879
misc_feature
/note="assembly_name:Contig8"
3980..6597
misc_feature
/note="assembly_name:Contig9"
6598..10040
misc_feature
/note="assembly_name:Contig10"
clone_end:SP6
vector_side:left
10141..12973
misc_feature
/note="assembly_name:Contig11"
13074..15987
misc_feature
/note="assembly_name:Contig12"
16088..19014
misc_feature
/note="assembly_name:Contig13"
19115..22164
misc_feature
/note="assembly_name:Contig14"
22265..26798
misc_feature
/note="assembly_name:Contig15"
26899..34198
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/note="assembly_name:Contig16"
34299..42840
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/note="assembly_name:Contig17"
42941..48833
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/note="assembly_name:Contig18"
48934..59689
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/note="assembly_name:Contig19"
59790..72021
misc_feature
/note="assembly_name:Contig20"
72122..87123
misc_feature
/note="assembly_name:Contig21"
87224..106312
misc_feature
/note="assembly_name:Contig22"
106413..133985
misc_feature
/note="assembly_name:Contig23"
134086..159723
misc_feature
/note="assembly_name:Contig24"
1728 others

BASE COUNT 48651 a 29918 c 29977 g 49449 t 1728 others
ORIGIN

Query Match 3.8%; Score 23; DB 66; Length 159723;
Best Local Similarity 100.0%; Pred. No. 0.078;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 163 agaattatctagccccaaatgtc 185
|||||
Db 3629 AGAATTATCTAGCCCCAAATGTC 3651

RESULT 15
AC078953 171589 bp DNA HTG 07-JAN-2001
LOCUS Homo sapiens chromosome 3q clone RP11-370M22, *** SEQUENCING IN
DEFINITION PROGRESS ***, 61 unordered pieces.
ACCESSION AC078953
VERSION AC078953.12 GI:12039127

Wed Nov 7 09:21:18 2001

HTG: HTGS_PHASE1.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE
AUTHORS

1 (bases 1 to 171589)
Muzny, D.M., Adams, C., Adio-Oduola, B., Ali-Osman, F.R., Allen, C.,
Alsbrooks, S.L., Amaratunga, H.C., Are, J.R., Banks, T., Barbaria, J.,
Benton, J., Blum, K., Blankenburg, K., Bonnin, D., Bouck, J.,
Bowie, S., Brieve, M., Brown, E., Brown, M., Bryant, N.P., Buhay, C.,
Burch, P., Burkett, C., Burrell, K.L., Byrd, N.C., Carron, T.F.,
Carter, M., Cavazos, S.R., Chacko, J., Chavez, D., Chen, G., Chen, R.,
Chen, Z., Chowdhry, I., Christopoulos, C., Cleveland, C.D., Cox, C.,
Coyle, M.D., Dathorne, R., David, R., Davila, M.L., Davis, C.,
Davy-Carroll, L., Dederich, D.A., Delaney, K.R., Delgado, O.,
Denn, A.L., Ding, Y., Dinh, H.H., Douthwaite, K.J., Draper, H.,
Dugan-Rocha, S., Durbin, K.J., Earnhart, C., Edgar, D., Edwards, C.C.,
Elhaj, C., Escotto, M., Falls, T., Ferraguto, D., Flagg, N., Ford, J.,
Foster, P., Frantz, P., Gabisi, A., Gao, J., Garcia, A., Garner, T.,
Garza, N., Gill, R., Correll, J.H., Guevara, W., Gunaratne, P., Hale, S.,
Hamilton, K., Hernandez, C., Harris, K., Hart, M., Havlak, P., Hawes, A.,
Hernandez, J., Hernandez, O., Hodgson, A., Hogues, M., Holloway, C.,
Hollins, B., Homs, F., Howard, S., Huber, J., Johnson, R., Jolivet, S.,
Jackson, L.E., Jacobson, B., Jia, Y., Johnson, R., Jolivet, S.,
Joudah, S., Karlsson, E., Kelly, S., Khan, U., King, L., Korvah, J.,
Kovar, C., Kratovic, J., Kureshi, A., Landry, N., Leal, B., Lewis, L.C.,
Lewis, L., Li, J., Li, Z., Lichtarge, O., Lieu, C., Liu, J., Liu, W.,
Loulsegh, H., Lozano, R.J., Lu, X., Lucier, A., Lucier, R., Luna, R.,
Ma, J., Maheshwari, M., Mapua, P., Martin, R., Martindale, A.,
Martinez, E., Massey, E., Mawhney, E., McLeod, M.P., Meador, M.,
Mei, G., Metzger, M., Miner, G., Miner, Z., Mitchell, T., Mohabbat, K.,
Morgan, M., Morris, S., Moser, M., Neal, D., Newton, J., Newton, N.,
Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokenkwo, S.,
Ogih, M., Okwundu, G., Oragunye, N., Oviedo, R., Pace, A., Payton, B.,
Peery, J., Perez, L., Peters, L., Picketts, R., Primus, E., Pu, L.L.,
Quiles, M., Ren, Y., Rives, M., Rojas, A., Rojebokan, I., Rolfe, M.,
Ruiz, S., Savery, G., Scherer, S., Scott, G., Shen, H., Shoohtari, N.,
Slisdon, I., Sodergren, E., Sonaike, T., Sparks, A., Stanley, H.,
Stone, H., Sutton, A., Svatek, A., Tabor, P., Tamerisa, A., Tamerisa, K.,
Tang, H., Tansey, J., Taylor, C., Taylor, P., Telford, B., Thomas, N.,
Thomas, S., Umani, K., Vasquez, L., Vera, V., Villalón, D., Vinson, R.,
Wall, R., Wang, S., Ward-Moore, S., Warren, R., Washington, C.,
Watlington, S., Williams, G., Williamson, A., Wleczyk, R., Wooden, S.,
Worley, K., Wu, C., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Nelson, D.,
and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 171589)
Worley, K.C.

Direct Submission
Submitted (13-AUG-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Jan 5, 2001 this sequence version replaced gi:11995537.

Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu

Project information

Center project name: HBS2
Center clone name: RP11-370M22

Summary Statistics

Sequencing vector: M13, L08821
Chemistry: Dye-terminator Big Dye 48x of reads
Assembly: Dye-terminator Big Dye 48x of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 135345 bases at least Q40
Consensus quality: 149003 bases at least Q30
Consensus quality: 157760 bases at least Q20
Estimated insert size: 151105; sum-of-contigs estimation
Quality coverage: 0x in Q20 bases; agarose-gel estimation
Quality coverage: 2.5x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 61 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 9607: contig of 9607 bp in length
9608 9707: gap of unknown length
15420: contig of 5713 bp in length
15520: gap of unknown length
21934: contig of 6414 bp in length
21935 21934: gap of unknown length
28036: contig of 6062 bp in length
28037 28036: gap of unknown length
31973: contig of 5677 bp in length
31974 31973: gap of unknown length
33973: contig of 5443 bp in length
33974 33973: gap of unknown length
35117: contig of 4833 bp in length
35118 35117: gap of unknown length
44350: contig of 4215 bp in length
44351 44350: gap of unknown length
48664: gap of unknown length
48665 48664: contig of 4910 bp in length
48765 48765: gap of unknown length
53675 53674: gap of unknown length
53676 53675: contig of 4877 bp in length
58552 58551: gap of unknown length
58553 58552: contig of 4249 bp in length
63001: gap of unknown length
63002 63001: contig of 3583 bp in length
66683: gap of unknown length
66684 66683: contig of 2922 bp in length
69705 69705: gap of unknown length
69805 69805: contig of 3354 bp in length
73159 73159: gap of unknown length
73259 73259: gap of unknown length
76844: contig of 3585 bp in length
76845 76844: gap of unknown length
80240: contig of 3296 bp in length
80340 80340: gap of unknown length
83170 83170: contig of 2830 bp in length
83271 83270: gap of unknown length
85770 85770: contig of 2500 bp in length
85771 85770: gap of unknown length
88260 88260: contig of 2390 bp in length
88261 88260: gap of unknown length
91251 91251: contig of 2891 bp in length
91252 91251: gap of unknown length
91351 91351: contig of 3787 bp in length
95138 95138: gap of unknown length
95139 95139: contig of 2276 bp in length
95239 95239: gap of unknown length
97515 97515: gap of unknown length
97614 97614: contig of 2169 bp in length
99783 99783: gap of unknown length
99784 99783: contig of 2616 bp in length
102499 102499: gap of unknown length
102599 102599: contig of 2419 bp in length
105018 105018: gap of unknown length
105019 105019: contig of 2608 bp in length
107727 107727: gap of unknown length
107728 107727: contig of 3486 bp in length
111312 111312: contig of 3408 bp in length
111412 111412: gap of unknown length
114820 114820: contig of 3408 bp in length
114920 114920: gap of unknown length
117190 117190: contig of 2270 bp in length
117191 117191: gap of unknown length
117291 117291: contig of 1987 bp in length
117292 117291: gap of unknown length
119377 119377: gap of unknown length
119378 119377: contig of 1583 bp in length
120960 120960: contig of 1583 bp in length
121060 121060: gap of unknown length

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*	121061	122181:	contig	of 1121 bp	in length
*	122182	122281:	gap of	unknown length	
*	122882	124038:	contig	of 1757 bp	in length
*	124039	124138:	gap of	unknown length	
*	124139	125666:	contig	of 1528 bp	in length
*	125667	125766:	gap of	unknown length	
*	125767	127725:	contig	of 1959 bp	in length
*	127726	127825:	gap of	unknown length	
*	127826	130376:	contig	of 2551 bp	in length
*	130377	130476:	gap of	unknown length	
*	130477	132820:	contig	of 2344 bp	in length
*	132821	132920:	gap of	unknown length	
*	132921	135499:	contig	of 2579 bp	in length
*	135500	135599:	gap of	unknown length	
*	135600	137202:	contig	of 1603 bp	in length
*	137203	137302:	gap of	unknown length	
*	137303	139112:	contig	of 1810 bp	in length
*	139113	139212:	gap of	unknown length	
*	139213	140413:	contig	of 1201 bp	in length
*	140414	140513:	gap of	unknown length	
*	140514	142609:	contig	of 2036 bp	in length
*	142610	142709:	gap of	unknown length	
*	142710	144637:	contig	of 1928 bp	in length
*	144638	144737:	gap of	unknown length	
*	144738	146091:	contig	of 1354 bp	in length
*	146092	146191:	gap of	unknown length	
*	146192	147858:	contig	of 1667 bp	in length
*	147859	147958:	gap of	unknown length	
*	147959	149631:	contig	of 1673 bp	in length
*	149632	149731:	gap of	unknown length	
*	149732	150740:	contig	of 1009 bp	in length
*	150741	150840:	gap of	unknown length	
*	150841	152348:	contig	of 1508 bp	in length
*	152349	152448:	gap of	unknown length	
*	152449	153976:	contig	of 1528 bp	in length
*	153977	154076:	gap of	unknown length	
*	154077	155379:	contig	of 1303 bp	in length
*	155380	155479:	gap of	unknown length	
*	155480	157688:	contig	of 2209 bp	in length
*	157689	157788:	gap of	unknown length	
*	157789	159085:	contig	of 1297 bp	in length
*	159086	159185:	gap of	unknown length	
*	159186	160288:	contig	of 1103 bp	in length
*	160289	160388:	gap of	unknown length	
*	160389	162315:	contig	of 1927 bp	in length
*	162316	162415:	gap of	unknown length	
*	162416	165343:	contig	of 1128 bp	in length
*	165344	166473:	gap of	unknown length	
*	166474	167773:	contig	of 1130 bp	in length
*	167774	168364:	contig	of 1130 bp	in length

Query Match	3.8%	Score 23;	DB 75;	Length 171589;
Best Local Similarity	100.0%;	Pred. No. 0.079;		
Mismatches 23;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

Qy 163 agaattatctagccccaaatgtc 185
 |||||
 Db 1643 AGAATTATCTAGCCCCAAATGTC 1665

Search completed: November 5, 2001, 23:44:59
Job time: 2829 sec

us-09-656-668-198.rng

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model
Run on: November 5, 2001, 22:33:25 ; Search time 126.12 Seconds
(without alignments)
3017.034 Million cell updates/sec

Title: US-09-656-668-198
Perfect score: 606
Sequence: 1 tgaatttgcccttaccc.....aagctgtttgtgtcgcac 606

Scoring table: IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Searched: 730101 seqs, 313950809 residues 1460202
Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_0601.*
1: /SIDSL1/gcgdata/geneseq/geneseq/NA1980.DAT.*
2: /SIDSL1/gcgdata/geneseq/geneseq/NA1981.DAT.*
3: /SIDSL1/gcgdata/geneseq/geneseq/NA1982.DAT.*
4: /SIDSL1/gcgdata/geneseq/geneseq/NA1983.DAT.*
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22: /SIDSL1/gcgdata/geneseq/geneseq/NA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Match %	Score	Length	ID	Description
1	606	100.0	606	22	AAF95006 Human ovarian canc
2	82.4	13.6	80240	20	AAV83940 NC-contig derived
3	82.4	13.6	80595	20	AAV83939 HC-contig derived
4	73	12.0	2099	21	AAA27046 Human cell surface
5	73	12.0	2116	21	AAAC59845 Human secreted pro
6	71	11.7	2061	20	AAZ33929 Human PRO329 nucle
7	71	11.7	2061	21	AAZ33929 Human PRO329 (UNC2
8	68.2	11.3	208	21	AAAF78471 Human prostate can
9	67.4	11.1	1829	21	AAAF15763 Human secreted pro
10	66.2	10.9	119950	20	AAAX90201 Human yes1 gene.
11	65.8	10.9	3280	21	AAA26367 Human secreted pro

12	65.8	10.9	3300	21	AAA26449 Human secreted pro
13	64.8	10.7	7505	20	AAV83949 Bacterial artifi
14	64.6	10.7	1583	21	AAV79962 Human secreted pro
15	63.4	10.5	2020	21	AAAC93321 Human neuroblastom
16	61.2	10.1	1119	22	AAAF97864 Human kinsin-like
17	61.2	10.1	121162	21	AAAC66548 Human KIK-L2 gene.
18	60.6	10.0	11570	21	AAAC95905 Sequence of a cDNA
19	60.4	10.0	6314	14	AAQ40341 Nucleotide sequenc
20	60	9.9	50000	21	AAAF41139 Homo sapiens 20q13
21	56.2	9.3	10282	19	AAV09023 Retinoblastoma bin
22	56.2	9.3	162450	21	AAZ86967 Human biallelic po
23	56	9.2	251	19	AAAX10977 Human secreted pro
24	54.8	9.0	1920	21	AAAC59071 Human secreted pro
25	54.8	9.0	1921	22	AAAF32739 Human secreted pro
26	54.8	9.0	2499	21	AAAC80607 Human secreted pro
27	54.8	9.0	2522	21	AAAC80618 Lung cancer associ
28	54.8	9.0	2547	21	AAAF18213 Human ORFX ORF2097
29	54.6	9.0	6439	21	AAAC76542 Human KCMQ5 (KCM6q
30	54.2	8.9	125910	21	AAAC64370 Human flavin-conta
31	53.8	8.9	25464	19	AAAF57274 Human kidney amino
32	53.2	8.8	49998	20	AAAC23518 Human secreted pro
33	53	8.7	354	21	AAAC22185 Novel human polynu
34	52.8	8.7	381	22	AAAF66536 Human secreted pro
35	52.8	8.7	476	21	AAAC32147 Human secreted pro
36	52.8	8.7	693	20	AAAC16692 Human secreted pro
37	52.6	8.7	155	21	AAAC16692 Human secreted pro
38	52	8.6	438	20	AAAX97944 Human secreted pro
39	52	8.6	2423	21	AAAC98795 Human pancreatic c
40	51.6	8.5	936	22	AAAF58252 Oligonucleotide D1
41	51.6	8.5	936	22	AAAF58252 Oligonucleotide D1
42	51.6	8.5	936	22	AAAF58257 Oligonucleotide D2
43	51.6	8.5	936	22	AAAF58259 Oligonucleotide D2
44	51.6	8.5	936	22	AAAF58262 Oligonucleotide D1
45	51.6	8.5	938	22	AAAF58255 Oligonucleotide D1

ALIGNMENTS

RESULT 1
AAF95006
ID AAF95006 standard; DNA; 606 BP.
XX
AC AAF95006;
XX
DT 23-MAY-2001 (first entry)
XX
DE Human ovarian cancer associated coding sequence SEQ ID NO: 198.
XX
KW Human, ovarian cancer; vaccine; gene therapy; carcinoma; ds.
XX
OS Homo sapiens.
XX
PN WO200118046-A2.
XX
PD 15-MAR-2001.
XX
PF 08-SEP-2000; 2000WO-US24827.
XX
PR 10-SEP-1999; 99US-0394374.
PR 01-MAY-2000; 2000US-0561778.
PR 15-AUG-2000; 2000US-0640173.
PR 07-SEP-2000; 2000US-0656668.
XX
PA (CORI-) CORIXA CORP.
XX
PI Xu J, Stolk JA;
XX
DR WFI; 2001-211395/21.
XX
PT Isolated polypeptides associated with ovarian carcinomas, and the
PT nucleic acids that encode them, useful for the prevention diagnosis and
PT treatment of ovarian cancers -

QY	192	actgctgttgagaaacccacgcagcagcttactgctgagcgttcctacgtgagcgttgcctt	251
Db	66908	gccaaagtggagaaacctcattcttgcgtctcttcccttctacgttctcaatcaactgtt	66967
QY	252	gttctggcttctgtag	267
Db	66968	gttcttcagcattag	66983
RESULT	4		
AAA27046			
ID	AAA27046	standard; CDNA; 2099 BP.	
XX	AA27046;		
XX	22-AUG-2000	(first entry)	
XX	Human cell surface receptor protein cDNA sequence #3.		
DE	Human; HCSRp; cytostatic; antiarthritic; antirheumatic; antithrombotic; antiparasitic; immunosuppressive; antiarteriosclerotic; antibacterial; antiparasitic; neuroprotective; nontropic; anticonvulsant; cancer; leukaemia; melanoma; rheumatoid arthritis; asthma; atherosclerosis; akathesia; Alzheimer's diseases; multiple sclerosis; epilepsy; ss.		
XX	Homo sapiens.		
OS			
XX	Location/Qualifiers		
XX	Key	43..1119	
XX	CDS	/tag=a *HCSRp-3*	
XX		/product= *HCSRp-3*	
XX	W0200028032-A2.		
XX	18-MAY-2000.		
XX	12-NOV-1999;	99WO-US26742.	
XX	12-NOV-1998;	98US-0191280.	
XX	07-DEC-1998;	98US-0206647.	
XX	08-MAR-1999;	99US-0123404.	
XX	(INCY-) INCYTE PHARM INC.		
XX	Tang YT, Corley NC, Guegler KJ, Yue H, Baughn MR, Lal P;		
XX	Hillman JL, Bandman O, Azimzai Y, Au-Young J;		
XX	WPI; 2000-376546/32.		
XX	P-PSDB; AAY94336.		
XX	New human cell surface receptor protein and polynucleotide useful for diagnosis, prevention and treatment of cancer, immune disorders, infection and neuronal disorders		
XX	Claim 9; Page 89-90; 97pp; English.		
XX	The present sequence encodes a novel human cell surface receptor protein (HCSRp) designated HCSRp-3. The nucleotide sequence was identified in Incyte Clone 53181 from the CDNA library LNOONNOT02, which was made from RNA isolated from lymph node tissue. A number of Incyte Clones were used to assemble the consensus sequence. BLAST analysis showed that the sequence is homologous to IgG Fc receptor g583604. HCSRp and its antagonist are useful for preventing treating disorders associated with decreased or increased expression activity of HCSRp. Such disorders include cancers such as leukaemia and melanoma, immune disorders such as rheumatoid arthritis, asthma and atherosclerosis, bacterial and parasitic infections and neuronal disorders such as akathesia, Alzheimer's disease, multiple sclerosis epilepsy. Polynucleotides encoding HCSRps may be used as hybridisation probes to diagnose these conditions. Anti-HCSRp antibodies may be used as antagonists, as a targeting or delivery mechanism for bringing pharmaceutical agents into contact with cells or tissues expressing		

AAV83939
ID AAV83939 standard; DNA: 80595 BP.
XX AC
XX AAV83939;
XX AC
XX 03-MAR-1999 (first entry)
XX AC
XX HC-contig derived from normal human chromosome 10q25.2 region.
XX DE
XX Yeast artificial chromosome; YAC; probe: eukaryotic chromosome;
XX KW neocentromere; replication; extra-chromosomal element; segregation;
XX KW cell division; artificial chromosome; gene therapy; mardel(10);
XX KW human artificial chromosome; transgenic; chromosome 10; 10q25.2; ss.
XX KW
XX Homo sapiens.
XX OS
XX W09851790-A1.
XX PN
XX 19-NOV-1998.
XX PD
XX 13-MAY-1998; 98WO-AU00352.
XX PF
XX 26-AUG-1997; 97AU-0008791.
XX PR
XX 13-MAY-1997; 97AU-0006784.
XX PR
XX (AMRA-) AMRAD OPERATIONS PTY LTD.
XX PA
XX Cancellia MR, Choo K, Du Sart D;
XX PI
XX WPT; 1999-009773/01.
XX DR
XX New isolated nucleic acid comprising neocentromere sequences from
XX PT eukaryotic chromosome - used to produce replicable, segregating
XX PT artificial chromosomes that can carry large amounts of DNA for gene
XX PT therapy
XX PT
XX Claim 8; Fig 6; 540pp; English.
XX PS
XX The present sequence represents the HC-contig derived from normal human
XX CC chromosome 10, 10q25.2 region. This region can be naturally mutated to
XX CC produce an unusual chromosomal marker referred to as mardel(10). The
XX CC mardel(10) marker is mitotically stable and contains a functional
XX CC neocentromere at a location regarded as non-centromeric. This
XX CC neocentromere maps to q25.2 on chromosome 10. The specification describes
XX CC nucleic acid sequences derived from a eukaryotic chromosome, including a
XX CC neocentromere or its functional derivative or hybrid, that are able, in
XX CC a compatible cell, of replicating, acting as extra-chromosomal element
XX CC and segregating during cell division. The sequences can be used to
XX CC construct artificial chromosomes for use in gene therapy comprising a
XX CC replicable, segregating nucleic acid that confers a specific phenotype
XX CC on cells. Human artificial chromosomes can propagate in human cells and
XX CC carry large amounts of DNA (e.g. therapeutic genes), and, being
XX CC extra-chromosomal, they are not mutagenic. The artificial chromosomes
XX CC are also useful for generation of transgenic plants and animals, in
XX CC production of proteins and to make diagnostic reagents, e.g. for
XX CC expression of cytokines, receptors and growth factors, or to increase
XX CC the copy number of a gene in a cell. The constructs may also be
XX CC used for functional and structural analysis of chromosomes.
XX CC
XX Sequence 80595 BP: 23183 A; 16613 C; 16824 G; 23975 T; 0 other;
XX SQ

Query Match 13.6%; Score 82.4; DB 20; Length 80595;
Best Local Similarity 63.9%; Pred. No. 1.2e-15;
Matches 125; Conservative 0; Mismatches 71; Indels 0; Gaps 131

QY 72 tgggtgggaacatcgtactggcatctaatcatagagggcagtaagtctgtctaacaatc 131
||||| ||||||| ||||||| || ||| ||| ||||||| |||||||
Db 66788 tgaagtgggtgggtgctactggcatctagtggtgtgagaccagagatgctttaaacatc 66847

QY 132 tttaacagcacagacagagcccaacaaagagaattatctagcccaaatgtccataac 191
||||| ||||||| ||||||| ||| ||| ||||||| ||||||| |||
Db 66848 ccgcaagcacagagcagtcgccgacaaagaattatcttggcccaataatcagtg 66907

us-09-656-668-198.rng

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```
CC HCSR and for diagnosis of HCSR-related disorders. HCSR and its
CC catalytic or immunogenic fragments are useful for drug screening using
CC libraries of compounds.
XX Sequence 2099 BP; 528 A; 534 C; 501 G; 536 T; 0 other;
SQ

Query Match 12.0%; Score 73; DB 21; Length 2099;
Best Local Similarity 72.9%; Pred. No. 2e-13;
Matches 94; Conservative 0; Mismatches 35; Indels 0; Gaps 0;

Qy 85 tgctactgcatctaatgcatgagggcagtaagtgtctgaacatctttcaacgcacag 144
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1523 tgctactgcatctaatgcatgagggcagtaagtgtctgaacatcttaataatgcacag 1582
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

Qy 145 gacagagcccaaaaagaattatctagcccaaaatgtccataacactgtgttgaga 204
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1583 ggcagtaccaccaagaagaataatctgcccacaaatgtcagtgtactgagttgaga 1642
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

Qy 205 aacactacc 213
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1643 aacccacgc 1651

RESULT 5
AAC59845
ID AAC59845 standard; DNA: 2116 BP.
XX AAC59845;
AC AAC59845;
XX
DT
DT
XX 26-JAN-2001 (first entry)
XX
DE Human secreted protein encoding DNA clone vq24 1.
XX
XX Secreted protein; human; autoimmune disorder; multiple sclerosis; ulcer;
KW systemic lupus erythematosus; rheumatoid arthritis; anaemia; stroke;
KW haematopoiesis regulation; tissue regrowth; wound healing; haemophilia;
KW Alzheimer's disease; Parkinson's disease; Shy-drager syndrome; cancer;
KW contraceptive; infection; growth inhibition; hyperproliferative disorder;
KW psoriasis; ds.
XX
XX Homo sapiens.
OS
OS WO200055375-A1.
PN
XX
XX 21-SEP-2000.
PD
XX
XX 17-MAR-2000; 2000WO-US07285.
PF
XX
XX 17-MAR-1999; 99US-0124809.
PR
XX 17-MAR-1999; 99US-0124916.
PR
XX 17-AUG-1999; 99US-0149639.
PR
XX 01-OCT-1999; 99US-0157247.
PR
XX 29-NOV-1999; 99US-0167824.
PR
XX 15-FEB-2000; 2000US-0182711.
XX
XX (ALPH-) ALPHAGENE INC.
PA
XX
XX Valenzuela D, Yuan O, Hoffman H, Hall J, Raple'jko P;
PI
XX
XX WPI; 2000-638211/61.
DR
XX P-PSDB; AAB34744.
DR
XX
XX Novel proteins and polypeptides useful for the treatment of e.g
PT multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis,
PT cancer, Alzheimer's disease, Parkinson's disease, stroke, anaemia and
PT ulcers.
PT
XX
XX Claim 124; Page 462; 493pp; English.
PS
XX
XX This invention relates to 59 human secreted proteins and the nucleotide
XX sequences encoding them. Sequences AAC59845-59846 and AAB34687-B34745
XX represent the proteins and their encoding nucleotide sequences, and
CC
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CC sequences AAB34746-B34771 represent fragments of the proteins. Probes
CC for the DNA sequences are represented by sequences AAC59847-C59596. The
CC proteins exhibit neuroprotective, dermatological, immunosuppressive,
CC antiinflammatory, antianemic, nootropic, antiparkinsonian,
CC cerebroprotective, haemostatic, vulnerary, cytostatic, antipsoriatic,
CC antibacterial, virucide, and fungicide activity. The proteins and
CC nucleotide sequences are useful as nutritional sources or supplements
CC and in research. The proteins are useful for treating immune deficiency
CC and disorders, which may be genetic or resulting from infections,
CC autoimmune disorders such as multiple sclerosis, systemic lupus
CC erythematosus, rheumatoid arthritis, and for treating myeloid or lymphoid
CC cell deficiencies such as anaemias by regulating haematopoiesis. The
CC proteins are also useful in compositions for bone, cartilage, tendon,
CC ligament and/or nerve tissue growth or regeneration, for wound healing,
CC tissue repair and replacement and in the treatment of wounds, incisions
CC and ulcers. Other uses include in the treatment of central and
CC peripheral nervous system and neuropathies such as Alzheimer's and
CC Parkinson's diseases and Shy-Drager syndrome, and mechanical and
CC traumatic disorders, such as spinal cord disorders, head trauma and
CC stroke. The proteins may also be used as a contraceptive, and for
CC treating coagulation disorders such as haemophilias. The protein and
CC nucleotide sequences with cadherin include for inhibiting the growth,
CC cancer. Other uses for the protein include for infecting agents such as bacteria,
CC infection or function of, or killing, infecting bodily characteristics
CC virus, fungi and other parasites, for effecting biorhythms or cardiac
CC cycles or rhythms, effecting metabolism, catabolism, anabolism,
CC processing, utilization, storage or elimination of dietary fat, lipid,
CC protein, carbohydrate, vitamins, minerals, cofactors, effecting
CC behavioural characteristics, providing analgesic effects and for treating
CC hyperproliferative disorders such as psoriasis.
XX Sequence 2116 BP; 545 A; 531 C; 503 G; 537 T; 0 other;
SQ

Query Match 12.0%; Score 73; DB 21; Length 2116;
Best Local Similarity 72.9%; Pred. No. 2e-13;
Matches 94; Conservative 0; Mismatches 35; Indels 0; Gaps 0;

Qy 85 tgctactgcatctaatgcatgagggcagtaagtgtctgaacatctttcaacgcacag 144
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1520 tgctactgcatctaatgcatgagggcagtaagtgtctgaacatcttaataatgcacag 1579
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

Qy 145 gacagagcccaaaaagaattatctagcccaaaatgtccataacactgtgttgaga 204
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1580 ggcagtaccaccaagaagaataatctgcccacaaatgtcagtgtactgagttgaga 1639
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

Qy 205 aacactacc 213
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1640 aacccacgc 1648

RESULT 6
AAC33929
ID AAC33929 standard; cDNA: 2061 BP.
XX AAC33929;
AC AAC33929;
XX
XX 07-DEC-1999 (first entry)
DT
XX
XX Human PRO329 nucleotide sequence.
XX
XX Human; PRO; EST; expressed sequence tag; PCR primer; hybridisation;
KW probe; blood coagulation disorder; cancer; cellular adhesion disorder;
KW secreted protein; transmembrane protein; ss.
XX
XX Homo sapiens.
OS
XX WO946281-A2.
PN
XX
XX 16-SEP-1999.
PD
XX 08-MAR-1999; 99WO-US05028.
PF
```

XX	10-MAR-1998;	98US-0077450.
PR	11-MAR-1998;	98US-0077632.
PR	11-MAR-1998;	98US-0077641.
PR	11-MAR-1998;	98US-0077649.
PR	12-MAR-1998;	98US-0077791.
PR	13-MAR-1998;	98US-0078004.
PR	17-MAR-1998;	98US-0040220.
PR	17-MAR-1998;	98US-0078886.
PR	20-MAR-1998;	98US-0078910.
PR	20-MAR-1998;	98US-0078936.
PR	20-MAR-1998;	98US-0078939.
PR	25-MAR-1998;	98US-0079294.
PR	25-MAR-1998;	98US-0079656.
PR	27-MAR-1998;	98US-0079663.
PR	27-MAR-1998;	98US-0079664.
PR	27-MAR-1998;	98US-0079689.
PR	27-MAR-1998;	98US-0079728.
PR	27-MAR-1998;	98US-0079786.
PR	30-MAR-1998;	98US-0079920.
PR	30-MAR-1998;	98US-0079923.
PR	31-MAR-1998;	98US-0080105.
PR	31-MAR-1998;	98US-0080107.
PR	31-MAR-1998;	98US-0080165.
PR	31-MAR-1998;	98US-0080324.
PR	01-APR-1998;	98US-0080327.
PR	01-APR-1998;	98US-0080328.
PR	01-APR-1998;	98US-0080333.
PR	01-APR-1998;	98US-0080334.
PR	08-APR-1998;	98US-00801049.
PR	08-APR-1998;	98US-00801070.
PR	08-APR-1998;	98US-00801071.
PR	09-APR-1998;	98US-0081195.
PR	09-APR-1998;	98US-0081203.
PR	09-APR-1998;	98US-0081229.
PR	15-APR-1998;	98US-0081817.
PR	15-APR-1998;	98US-0081838.
PR	15-APR-1998;	98US-0081952.
PR	15-APR-1998;	98US-0081955.
PR	21-APR-1998;	98US-0082568.
PR	21-APR-1998;	98US-0082569.
PR	22-APR-1998;	98US-0082700.
PR	22-APR-1998;	98US-0082704.
PR	22-APR-1998;	98US-0082767.
PR	23-APR-1998;	98US-0082796.
PR	23-APR-1998;	98US-0083336.
PR	27-APR-1998;	98US-0083322.
PR	28-APR-1998;	98US-0083392.
PR	29-APR-1998;	98US-0083495.
PR	29-APR-1998;	98US-0083496.
PR	29-APR-1998;	98US-0083499.
PR	29-APR-1998;	98US-0083500.
PR	29-APR-1998;	98US-0083545.
PR	29-APR-1998;	98US-0083554.
PR	29-APR-1998;	98US-0083558.
PR	29-APR-1998;	98US-0083559.
PR	30-APR-1998;	98US-0083742.
PR	05-MAY-1998;	98US-0084366.
PR	06-MAY-1998;	98US-0084414.
PR	06-MAY-1998;	98US-0084441.
PR	07-MAY-1998;	98US-0084450.
PR	07-MAY-1998;	98US-0084600.
PR	07-MAY-1998;	98US-0084627.
PR	07-MAY-1998;	98US-0084637.
PR	07-MAY-1998;	98US-0084639.
PR	07-MAY-1998;	98US-0084640.
PR	13-MAY-1998;	98US-0084643.
PR	13-MAY-1998;	98US-0085323.
PR	13-MAY-1998;	98US-0085338.
PR	13-MAY-1998;	98US-0085339.
PR	15-MAY-1998;	98US-0085573.
PR	15-MAY-1998;	98US-0085579.

15-MAY-1998: 98US-0085580.
PR 98US-0085582.
PR 15-MAY-1998: 98US-0085689.
PR 15-MAY-1998: 98US-0085697.
PR 15-MAY-1998: 98US-0085907.
PR 15-MAY-1998: 98US-0085704.
PR 15-MAY-1998: 98US-0086023.
PR 18-MAY-1998: 98US-0086392.
PR 22-MAY-1998: 98US-0086414.
PR 22-MAY-1998: 98US-0086430.
PR 22-MAY-1998: 98US-0086486.
PR 22-MAY-1998: 98US-0087098.
PR 28-MAY-1998: 98US-0087106.
PR 28-MAY-1998: 98US-0087208.
PR 28-MAY-1998: 98US-0087208.
PR 30-JUL-1998: 98US-0094651.
PR 11-SEP-1998: 98US-0100038.
XX
XX (GETH) GENENTECH INC.
PA
XX Wood WI, Goddard A, Gurney A, Yuan J, Baker KP, Chen J;
PI
XX WPI; 1999-551358/46.
XX P-PSDB; AA441690.
DR
XX New secreted and transmembrane polypeptides and their polynucleotides,
XX useful for treating blood coagulation disorders, cancers and cellular
PT adhesion disorders -
PT
PT
XX Claim 2; Fig 19; 530pp; English.
PS
XX The present invention describes secreted and transmembrane polypeptides
XX and their polynucleotides. The nucleotide sequences are useful as
CC sources of probes, primers, for chromosome mapping, and for generation
CC of antisense sequences. They can also be used to create transgenic
CC animals. The proteins can be used to treat a variety of diseases and
CC disorders, depending on their function. Diseases that may be treated
CC include blood coagulation disorders, cancers and cellular adhesion
CC disorders. They may also be used to raise antibodies. AA233891 to
CC AA234338, and AA41685 to AA41774 represent polynucleotide and
CC polypeptide sequence given in the exemplification of the present
CC invention.
XX
XX Sequence 2061 BP; 511 A; 528 C; 491 G; 531 T; 0 other;
SQ

[illegible]

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expressed sequence tag; detection; cancer; ss.

XX OS Homo sapiens.

XX PN WO200053756-A2.

XX PD 14-SEP-2000.

XX PF 18-FEB-2000; 2000WO-US04341.

XX PR 08-MAR-1999; 99WO-US05028.

XX PR 12-MAR-1999; 99US-0123957.

XX PR 29-MAR-1999; 99US-0126773.

XX PR 21-APR-1999; 99US-0130232.

XX PR 28-APR-1999; 99US-0131445.

XX PR 14-MAY-1999; 99US-0134287.

XX PR 23-JUN-1999; 99US-0141037.

XX PR 26-JUL-1999; 99US-0145698.

XX PR 29-OCT-1999; 99US-0162506.

XX PR 30-NOV-1999; 99WO-US28313.

XX PR 02-DEC-1999; 99WO-US28551.

XX PR 02-DEC-1999; 99WO-US28565.

XX PR 16-DEC-1999; 99WO-US30095.

XX PR 30-DEC-1999; 99WO-US31243.

XX PR 30-DEC-1999; 99WO-US31274.

XX PR 05-JAN-2000; 2000WO-US00219.

XX PR 06-JAN-2000; 2000WO-US00277.

XX PR 06-JAN-2000; 2000WO-US00376.

XX PA (GETH) GENENTECH INC.

XX PI Ashkenazi AJ, Baker KP, Botstein D, Desnoyers L, Eaton DL;

XX PI Ferrara N, Filvaroff E, Fong S, Gao W, Gerber H, Gerritsen ME;

XX PI Goddard A, Godowski PJ, Grimaldi CJ, Gurney AL, Hillan KJ;

XX PI Kljavin IJ, Kuo SS, Napier MA, Pan J, Paoni NF, Roy MA;

XX PI Shelton DL, Stewart TA, Tumas D, Williams PM, Wood WI;

XX DR WPI; 2000-611443/58.

XX DR P-PSDB; AAB44246.

XX PT Novel PRO polypeptides and polynucleotides used in detection methods,

XX PT to target bioactive molecules to specific cells, and to modulate

XX PT cellular activities -

XX PS Claim 2; Fig 19; 636pp; English.

XX CC AAC78458 to AAC78599 represent polynucleotide and EST (expressed

XX CC sequence tag) sequences which encode secreted or transmembrane PRO

XX CC polypeptides. The PRO polynucleotides and polypeptides have cytostatic

XX CC activity. The polynucleotides and polypeptides can be used for detecting

XX OS Homo sapiens.

XX PN WO200053756-A2.

XX PD 14-SEP-2000.

XX PF 18-FEB-2000; 2000WO-US04341.

XX PR 08-MAR-1999; 99WO-US05028.

XX PR 12-MAR-1999; 99US-0123957.

XX PR 29-MAR-1999; 99US-0126773.

XX PR 21-APR-1999; 99US-0130232.

XX PR 28-APR-1999; 99US-0131445.

XX PR 14-MAY-1999; 99US-0134287.

XX PR 23-JUN-1999; 99US-0141037.

XX PR 26-JUL-1999; 99US-0145698.

XX PR 29-OCT-1999; 99US-0162506.

XX PR 30-NOV-1999; 99WO-US28313.

XX PR 02-DEC-1999; 99WO-US28551.

XX PR 02-DEC-1999; 99WO-US28565.

XX PR 16-DEC-1999; 99WO-US30095.

XX PR 30-DEC-1999; 99WO-US31243.

XX PR 30-DEC-1999; 99WO-US31274.

XX PR 05-JAN-2000; 2000WO-US00219.

XX PR 06-JAN-2000; 2000WO-US00277.

XX PR 06-JAN-2000; 2000WO-US00376.

XX PA (GETH) GENENTECH INC.

XX PI Ashkenazi AJ, Baker KP, Botstein D, Desnoyers L, Eaton DL;

XX PI Ferrara N, Filvaroff E, Fong S, Gao W, Gerber H, Gerritsen ME;

XX PI Goddard A, Godowski PJ, Grimaldi CJ, Gurney AL, Hillan KJ;

XX PI Kljavin IJ, Kuo SS, Napier MA, Pan J, Paoni NF, Roy MA;

XX PI Shelton DL, Stewart TA, Tumas D, Williams PM, Wood WI;

XX DR WPI; 2000-611443/58.

XX DR P-PSDB; AAB44246.

XX PT Novel PRO polypeptides and polynucleotides used in detection methods,

XX PT to target bioactive molecules to specific cells, and to modulate

XX PT cellular activities -

XX PS Claim 2; Fig 19; 636pp; English.

XX CC AAC78458 to AAC78599 represent polynucleotide and EST (expressed

XX CC sequence tag) sequences which encode secreted or transmembrane PRO

XX CC polypeptides. The PRO polynucleotides and polypeptides have cytostatic

Qy 207 acctacc 213

Db 1613 cccagc 1619

RESULT 8

AAF15763

ID AAF15763 standard; cDNA; 208 BP.

XX AC AAF15763;

XX DT 13-MAR-2001 (first entry)

XX DE Human prostate cancer antigen nucleotide sequence SEQ ID NO:198.

XX KW Human; prostate cancer; prostate cancer antigen; detection; diagnosis;

XX KW neuroprotective; cytostatic; cardioactive; immunomodulatory; muscular;

XX KW vulnetary; gastrointestinal; nephrotropic; antinefactive; gynaecological;

XX KW antibacterial; gene therapy; neural; immune; reproductive; renal;

XX KW gastrointestinal; pulmonary; cardiovascular; proliferative disorder;

XX KW wound; infectious disease; ss.

XX OS Homo sapiens.

XX PN WO200055174-A1.

XX PD 21-SEP-2000.

XX PF 08-MAR-2000; 2000WO-US05988.

XX PR 12-MAR-1999; 99US-0124270.

XX PA (HUMA-) HUMAN GENOME SCI INC.

XX PA (ROSE/) ROSEN C A.

XX PI Rosen CA, Ruben SM;

XX PI WPI; 2000-587513/55.

XX DR P-PSDB; AAB56560.

XX PT Prostate cancer associated gene sequences, referred to as prostate

XX PT cancer antigens, useful for treatment, prevention, and diagnosis of

XX PT disorders such as prostate cancer -

XX PS Claim 1; Page 736; 2338pp; English.

XX CC AAF15566 to AAF16505 encode the human prostate cancer associated

XX CC proteins, called prostate cancer antigens, given in AAB56363 to AAB57302.

XX CC The prostate cancer antigens can have neuroprotective, cytostatic,

XX CC cardioactive, immunomodulatory, muscular, vulnetary, gastrointestinal,

XX CC nephrotropic, antinefactive, gynaecological and antibacterial activities,

XX CC and can be used in gene therapy. The prostate cancer antigen

XX CC polynucleotides may be used for detection of prostate cancer, chromosome

XX CC identification, as chromosome markers, and for numerous other diagnostic

XX CC or research purposes. The prostate cancer antigens may be used to treat

XX CC disorders such as neural, immune, muscular, reproductive, proliferative

XX CC gastrointestinal, pulmonary, cardiovascular, renal, and infectious

XX CC disorders, wounds, and infectious diseases. AAF16506 to AAF16514 to

XX CC AAB57303 represent sequences used in the exemplification of the present

XX CC invention.

XX SQ Sequence 208 BP; 63 A; 38 C; 54 G; 51 T; 2 other;

Query Match 11.7%; Score 71; DB 21; Length 2061;

Best Local Similarity 72.4%; Pred. No. 8.7e-13;

Matches 92; Conservative 0; Mismatches 35; Indels 0; Gaps 0;

Qy 87 ctactggcatcctaatgcagtagggcagtaatgtgttaaacatctttcaacgcacagga 146

Db 1493 ctactggcatcagtaataagaccagggtgcgcgttaaacatcctataatgcacagg 1552

Qy 147 cagagcccccacaaaggaattatctagcccaaatgtccataaacactgtgttgagaaa 206

Db 1553 cagtaccaccacacgaataatctgtgccccaaatgtcagttgtactgagtttgagaaa 1612

us-09-656-668-198.rng

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11.1%; Score 67.4; DB 21; Length 1829;
 Query Match 11.1%; Score 67.4; DB 21; Length 1829;
 Best Local Similarity 67.8%; Pred. No. 1.2e-11;
 Matches 139; Conservative 0; Mismatches 61; Indels 5; Gaps 3;
 21 catccagtgatattgcaattcctcaaaagacgtgtttttgattgtcacacacgtggtggg 80
 89 cctccagggagacagttgaaatgtctagagac--atttgattattatgctggcaggac 146
 81 acatgctactggcatctaatgcatagagggcagtaagtgtctaaacatcttcaacgc 140
 147 gcc--ctagtggcatctagtgtgagggtaagggtgctgtaagcgtctccataac 204
 141 acaggacagagcccccacaaagaaattctctagcccccaaatgcccataacactgtgt 200
 205 acaggacagacccccccacaa--agaattatccaccccccaaatgctagtagtgcagggt 263
 201 gagaaacactaccgaggatcttac 225
 264 gagaaacccactctgtctctaac 288
 RESULT 10
 AAX90201
 ID AAX90201 standard; DNA; 119950 BP.
 XX AAX90201;
 AC AAX90201;
 XX AAX90201;
 DT 23-SEP-1999 (first entry)
 XX Human yes1 gene.
 DE Human; yes1; diagnosis; neuropsychiatric disorder; BAD; schizophrenia;
 KW bipolar affective disorder; attention deficit disorder;
 KW schizoaffective disorder; unipolar affective disorder;
 KW Huntington's disease; Parkinson's disease; manic-depression; ds.
 XX Homo sapiens.
 OS Homo sapiens.
 XX WO9935290-A1.
 PN 15-JUL-1999.
 PD 07-JAN-1999; 99WO-US00297.
 PF 08-JAN-1998; 98US-0003944.
 PR (MILL-) MILLENNIUM PHARM INC.
 XX Chen H, Freimer NB;
 PI WPI: 1999-444203/37.
 XX P-PSDB; AAY24421.
 DR Detection of a genetic mutation in the yes1 gene, useful for
 XX diagnosis of a yes1 mediated neuropsychiatric disorder in a human
 PS Claim 1; Fig 2; 110pp; English.
 XX The present invention describes a method for detecting a genetic
 CC mutation in the yes1 gene for the diagnosis of a yes1 mediated
 CC neuropsychiatric disorder in a human. The method comprises detecting the
 CC presence or absence of a genetic mutation in the yes1 gene of the
 CC subject, where the genetic mutation is a substitution, insertion or a
 CC deletion and results in the production of a yes1 protein having an amino
 CC acid sequence other than the wild-type yes1 amino acid sequence and the
 CC presence of the genetic mutation identifies a subject that has or is at
 CC risk for developing a yes1 mediated neuropsychiatric disorder. Compounds
 CC that bind to the yes1 protein, alter the amount of the protein, or alter
 CC the activity of the yes1 gene product, are useful for treating a yes1
 CC mediated neuropsychiatric disorder. The disorders include Huntington's
 CC disease, Parkinson's disease, and especially bipolar-affective disorder

11.1%; Score 67.4; DB 21; Length 1829;
 Query Match 11.1%; Score 67.4; DB 21; Length 1829;
 Best Local Similarity 67.8%; Pred. No. 1.2e-11;
 Matches 139; Conservative 0; Mismatches 61; Indels 5; Gaps 3;
 21 catccagtgatattgcaattcctcaaaagacgtgtttttgattgtcacacacgtggtggg 80
 89 cctccagggagacagttgaaatgtctagagac--atttgattattatgctggcaggac 146
 81 acatgctactggcatctaatgcatagagggcagtaagtgtctaaacatcttcaacgc 140
 147 gcc--ctagtggcatctagtgtgagggtaagggtgctgtaagcgtctccataac 204
 141 acaggacagagcccccacaaagaaattctctagcccccaaatgcccataacactgtgt 200
 205 acaggacagacccccccacaa--agaattatccaccccccaaatgctagtagtgcagggt 263
 201 gagaaacactaccgaggatcttac 225
 264 gagaaacccactctgtctctaac 288
 RESULT 10
 AAX90201
 ID AAX90201 standard; DNA; 119950 BP.
 XX AAX90201;
 AC AAX90201;
 XX AAX90201;
 DT 23-SEP-1999 (first entry)
 XX Human yes1 gene.
 DE Human; yes1; diagnosis; neuropsychiatric disorder; BAD; schizophrenia;
 KW bipolar affective disorder; attention deficit disorder;
 KW schizoaffective disorder; unipolar affective disorder;
 KW Huntington's disease; Parkinson's disease; manic-depression; ds.
 XX Homo sapiens.
 OS Homo sapiens.
 XX WO9935290-A1.
 PN 15-JUL-1999.
 PD 07-JAN-1999; 99WO-US00297.
 PF 08-JAN-1998; 98US-0003944.
 PR (MILL-) MILLENNIUM PHARM INC.
 XX Chen H, Freimer NB;
 PI WPI: 1999-444203/37.
 XX P-PSDB; AAY24421.
 DR Detection of a genetic mutation in the yes1 gene, useful for
 XX diagnosis of a yes1 mediated neuropsychiatric disorder in a human
 PS Claim 1; Fig 2; 110pp; English.
 XX The present invention describes a method for detecting a genetic
 CC mutation in the yes1 gene for the diagnosis of a yes1 mediated
 CC neuropsychiatric disorder in a human. The method comprises detecting the
 CC presence or absence of a genetic mutation in the yes1 gene of the
 CC subject, where the genetic mutation is a substitution, insertion or a
 CC deletion and results in the production of a yes1 protein having an amino
 CC acid sequence other than the wild-type yes1 amino acid sequence and the
 CC presence of the genetic mutation identifies a subject that has or is at
 CC risk for developing a yes1 mediated neuropsychiatric disorder. Compounds
 CC that bind to the yes1 protein, alter the amount of the protein, or alter
 CC the activity of the yes1 gene product, are useful for treating a yes1
 CC mediated neuropsychiatric disorder. The disorders include Huntington's
 CC disease, Parkinson's disease, and especially bipolar-affective disorder

11.1%; Score 67.4; DB 21; Length 1829;
 Query Match 11.1%; Score 67.4; DB 21; Length 1829;
 Best Local Similarity 67.8%; Pred. No. 1.2e-11;
 Matches 139; Conservative 0; Mismatches 61; Indels 5; Gaps 3;
 21 catccagtgatattgcaattcctcaaaagacgtgtttttgattgtcacacacgtggtggg 80
 89 cctccagggagacagttgaaatgtctagagac--atttgattattatgctggcaggac 146
 81 acatgctactggcatctaatgcatagagggcagtaagtgtctaaacatcttcaacgc 140
 147 gcc--ctagtggcatctagtgtgagggtaagggtgctgtaagcgtctccataac 204
 141 acaggacagagcccccacaaagaaattctctagcccccaaatgcccataacactgtgt 200
 205 acaggacagacccccccacaa--agaattatccaccccccaaatgctagtagtgcagggt 263
 201 gagaaacactaccgaggatcttac 225
 264 gagaaacccactctgtctctaac 288
 RESULT 10
 AAX90201
 ID AAX90201 standard; DNA; 119950 BP.
 XX AAX90201;
 AC AAX90201;
 XX AAX90201;
 DT 23-SEP-1999 (first entry)
 XX Human yes1 gene.
 DE Human; yes1; diagnosis; neuropsychiatric disorder; BAD; schizophrenia;
 KW bipolar affective disorder; attention deficit disorder;
 KW schizoaffective disorder; unipolar affective disorder;
 KW Huntington's disease; Parkinson's disease; manic-depression; ds.
 XX Homo sapiens.
 OS Homo sapiens.
 XX WO9935290-A1.
 PN 15-JUL-1999.
 PD 07-JAN-1999; 99WO-US00297.
 PF 08-JAN-1998; 98US-0003944.
 PR (MILL-) MILLENNIUM PHARM INC.
 XX Chen H, Freimer NB;
 PI WPI: 1999-444203/37.
 XX P-PSDB; AAY24421.
 DR Detection of a genetic mutation in the yes1 gene, useful for
 XX diagnosis of a yes1 mediated neuropsychiatric disorder in a human
 PS Claim 1; Fig 2; 110pp; English.
 XX The present invention describes a method for detecting a genetic
 CC mutation in the yes1 gene for the diagnosis of a yes1 mediated
 CC neuropsychiatric disorder in a human. The method comprises detecting the
 CC presence or absence of a genetic mutation in the yes1 gene of the
 CC subject, where the genetic mutation is a substitution, insertion or a
 CC deletion and results in the production of a yes1 protein having an amino
 CC acid sequence other than the wild-type yes1 amino acid sequence and the
 CC presence of the genetic mutation identifies a subject that has or is at
 CC risk for developing a yes1 mediated neuropsychiatric disorder. Compounds
 CC that bind to the yes1 protein, alter the amount of the protein, or alter
 CC the activity of the yes1 gene product, are useful for treating a yes1
 CC mediated neuropsychiatric disorder. The disorders include Huntington's
 CC disease, Parkinson's disease, and especially bipolar-affective disorder

11.1%; Score 67.4; DB 21; Length 1829;
 Query Match 11.1%; Score 67.4; DB 21; Length 1829;
 Best Local Similarity 67.8%; Pred. No. 1.2e-11;
 Matches 139; Conservative 0; Mismatches 61; Indels 5; Gaps 3;
 21 catccagtgatattgcaattcctcaaaagacgtgtttttgattgtcacacacgtggtggg 80
 89 cctccagggagacagttgaaatgtctagagac--atttgattattatgctggcaggac 146
 81 acatgctactggcatctaatgcatagagggcagtaagtgtctaaacatcttcaacgc 140
 147 gcc--ctagtggcatctagtgtgagggtaagggtgctgtaagcgtctccataac 204
 141 acaggacagagcccccacaaagaaattctctagcccccaaatgcccataacactgtgt 200
 205 acaggacagacccccccacaa--agaattatccaccccccaaatgctagtagtgcagggt 263
 201 gagaaacactaccgaggatcttac 225
 264 gagaaacccactctgtctctaac 288
 RESULT 10
 AAX90201
 ID AAX90201 standard; DNA; 119950 BP.
 XX AAX90201;
 AC AAX90201;
 XX AAX90201;
 DT 23-SEP-1999 (first entry)
 XX Human yes1 gene.
 DE Human; yes1; diagnosis; neuropsychiatric disorder; BAD; schizophrenia;
 KW bipolar affective disorder; attention deficit disorder;
 KW schizoaffective disorder; unipolar affective disorder;
 KW Huntington's disease; Parkinson's disease; manic-depression; ds.
 XX Homo sapiens.
 OS Homo sapiens.
 XX WO9935290-A1.
 PN 15-JUL-1999.
 PD 07-JAN-1999; 99WO-US00297.
 PF 08-JAN-1998; 98US-0003944.
 PR (MILL-) MILLENNIUM PHARM INC.
 XX Chen H, Freimer NB;
 PI WPI: 1999-444203/37.
 XX P-PSDB; AAY24421.
 DR Detection of a genetic mutation in the yes1 gene, useful for
 XX diagnosis of a yes1 mediated neuropsychiatric disorder in a human
 PS Claim 1; Fig 2; 110pp; English.
 XX The present invention describes a method for detecting a genetic
 CC mutation in the yes1 gene for the diagnosis of a yes1 mediated
 CC neuropsychiatric disorder in a human. The method comprises detecting the
 CC presence or absence of a genetic mutation in the yes1 gene of the
 CC subject, where the genetic mutation is a substitution, insertion or a
 CC deletion and results in the production of a yes1 protein having an amino
 CC acid sequence other than the wild-type yes1 amino acid sequence and the
 CC presence of the genetic mutation identifies a subject that has or is at
 CC risk for developing a yes1 mediated neuropsychiatric disorder. Compounds
 CC that bind to the yes1 protein, alter the amount of the protein, or alter
 CC the activity of the yes1 gene product, are useful for treating a yes1
 CC mediated neuropsychiatric disorder. The disorders include Huntington's
 CC disease, Parkinson's disease, and especially bipolar-affective disorder

ID	AC79962 standard; cDNA; 1583 BP.
XX	AAC79962;
AC	12-FEB-2001 (first entry)
XX	Human secreted protein encoding cDNA for gene 15.
DE	Secreted protein; human; immunosuppressive; antiarthritic; antirheumatic; antiproliferative; cytostatic; cardiant; vasotropic; cerebroprotective; KW neurotropic; neuroprotective; antibacterial; virucide; fungicide; KW nontoxic; cerebrovascular disorder; cardiovascular disease; KW ophthalmological; vulnary; gene therapy; treatment; autoimmune disease; KW hyperproliferative disorder; cardiovascular disorder; ocular disorder; KW cerebrovascular disorder; nervous system disorder; infection; skin aging; KW wound healing; epithelial cell proliferation; transplantation; ss. XX Homo sapiens.
OS	WO200058357-A1.
AC	05-OCT-2000.
PX	23-MAR-2000; 2000WO-US07723.
PN	26-MAR-1999; 99US-0126506.
XX	07-JAN-2000; 2000US-0174852.
PR	(HUMA-) HUMAN GENOME SCI INC.
PD	Rosen CA, Ruben SM, Komatsoulis G;
PP	WPI: 2000-611704/58.
XX	P-PDSB: AAB45039.
DR	Nucleic acid molecules encoding human secreted proteins, used in preventing, treating or ameliorating a disorder, e.g. Alzheimer's and Parkinson's diseases and cancers -
PT	Claim 1a; Page 349; 418pp; English.
XX	This invention describes novel isolated nucleic acid molecules (I) encoding a human secreted proteins (II) which have immunosuppressive, antiarthritic, antirheumatic, antiproliferative, cytostatic, cardiant, vasotropic, cerebroprotective, nontoxic, neuroprotective, antibacterial, virucide, fungicide, ophthalmological and vulnary activity and can be used for gene therapy. (I) and (II) are used to prevent, treat or ameliorate a medical condition in e.g. humans, mice, rabbits, goats, horses, cats, dogs, chickens or sheep. (I) and (II) are also used in diagnosing a pathological condition or susceptibility to a pathological condition. The antibodies to (II) can also be used in alleviating symptoms associated with the disorders and in diagnostic immunoassays e.g. radioluminoassays or enzyme linked immunosorbent assays (ELISA). Disorders which are diagnosed or treated include autoimmune diseases e.g. rheumatoid arthritis, hyperproliferative disorders e.g. cardiac arrest, of the breast or liver, cardiovascular disorders e.g. cerebral ischemia, angiogenesis, nervous cerebrovascular disorders e.g. Alzheimer's disease, infections caused by bacterial system disorders e.g. viral diseases, corneal infection. The viruses and fungi and ocular disorders e.g. corneal infection. The polypeptides can also be used to aid wound healing and epithelial cell proliferation, to prevent skin aging due to sunburn, to maintain organs, before transplantation, for supporting cell culture of primary tissues, to regenerate tissues and in chemotaxis. The polypeptides can also be used as a food additive or preservative to increase or decrease storage capabilities.
CC	Sequence 1583 BP; 427 A; 370 C; 316 G; 458 T; 12 other;
XX	
SQ	

Query Match 10.7%; Score 64.6; DB 21; Length 1583;

Best Local Similarity 64.5%; Pred. No. 9.le-11;

Matches 91; Conservative 3; Mismatches 47; Indels 0; Gaps

Wed Nov 7 09:21:18 2001

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model
Run on: November 5, 2001, 23:00:40 ; Search time 126.05 Seconds
(without alignments)
3018.710 Million cell updates/sec

Title: US-09-656-668-198
Perfect score: 606
Sequence: 1 tgaagttgccccctaccce.....aagcctgtttgtctgtgcac 606

Scoring table: OLIGO_NVC
Gapop 60.0 , Gapext 60.0

Searched: 730101 seqs, 313950809 residues

Word size : 0
Total number of hits satisfying chosen parameters: 1460202

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : N_Geneseq_0601.*
1: /SIDSI/gcgdata/geneseq/geneseq/NA1980.DAT.*
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4: /SIDSI/gcgdata/geneseq/geneseq/NA1983.DAT.*
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12: /SIDSI/gcgdata/geneseq/geneseq/NA1991.DAT.*
13: /SIDSI/gcgdata/geneseq/geneseq/NA1992.DAT.*
14: /SIDSI/gcgdata/geneseq/geneseq/NA1993.DAT.*
15: /SIDSI/gcgdata/geneseq/geneseq/NA1994.DAT.*
16: /SIDSI/gcgdata/geneseq/geneseq/NA1995.DAT.*
17: /SIDSI/gcgdata/geneseq/geneseq/NA1996.DAT.*
18: /SIDSI/gcgdata/geneseq/geneseq/NA1997.DAT.*
19: /SIDSI/gcgdata/geneseq/geneseq/NA1998.DAT.*
20: /SIDSI/gcgdata/geneseq/geneseq/NA1999.DAT.*
21: /SIDSI/gcgdata/geneseq/geneseq/NA2000.DAT.*
22: /SIDSI/gcgdata/geneseq/geneseq/NA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	606	100.0	606	22 AAF95006	Human ovarian can
2	21	3.5	51	19 AAV36272	Probe used to isol
3	19	3.1	2769	21 AAC76385	Human ORFX ORF1940
4	18	3.0	1427	11 AAQ06902	Sequence encoding
5	18	3.0	1658	21 AAF21367	Human low adenosi
6	18	3.0	1658	21 AAA35245	Human adenosine re
7	18	3.0	1674	11 AAQ06901	Sequence encoding
8	18	3.0	2582	18 AAT87428	Clone J422. Homo
9	18	3.0	2635	19 AAV15587	Human B-cell activ
10	18	3.0	2697	20 AAV83816	cDNA encoding a B
11	18	3.0	2775	18 AAT72271	Human B cell surfa

c	12	18	3.0	2775	21	AAA08399	Human RPI05 nucleo
	13	18	3.0	3191	21	AAF21368	Human low adenosi
	14	18	3.0	3191	21	AAAC35246	Human adenosine re
	15	18	3.0	1830121	17	AAT42063	Haemophilus influe
	16	17	2.8	268	21	AAC33704	Arabidopsis thalia
	17	17	2.8	352	20	AAV86186	EST clone J638. H
	18	17	2.8	394	16	AAT26421	Human gene signatu
	19	17	2.8	396	20	AAV87210	EST clone BN99. H
	20	17	2.8	497	21	AAAC53869	Arabidopsis thalia
	21	17	2.8	542	21	AAAC53133	Arabidopsis thalia
	22	17	2.8	570	21	AAV65305	Human secreted pro
	23	17	2.8	776	18	AAT61702	Rat probasin encod
	24	17	2.8	1329	18	AAT67996	H. pylori cytoplas
	25	17	2.8	1383	18	AAT66164	Mouse interleukin-
	26	17	2.8	1551	21	AAC81117	Human secreted pro
	27	17	2.8	1680	22	AAF23411	Human SEC2 DNA seq
	28	17	2.8	2364	21	AAZ55926	cDNA encoding xeno
	29	17	2.8	2601	20	AAAX90316	Fervidobacterium p
	30	17	2.8	5921	20	AAZ32700	Human glycophospha
	31	17	2.8	8625	21	AAAF5834	Human ORFX ORF1389
	32	17	2.8	15512	19	AAV59470	Human ryanodin rec
	33	17	2.8	43795	21	AAZ92583	Human DAZ genomic
	34	17	2.8	240825	22	AAF24497	Human PG-3 gene.
	35	16	2.6	166	21	AAAC25754	Human secreted pro
	36	16	2.6	264	21	AAAC30995	Human secreted pro
	37	16	2.6	269	21	AAZ93797	Sequence encoding
	38	16	2.6	317	21	AAAC29682	Human secreted pro
	39	16	2.6	371	14	AAQ39834	Expressed Sequence
	40	16	2.6	371	14	AAQ59246	Human brain Expres
	41	16	2.6	375	21	AAAC26100	Human secreted pro
	42	16	2.6	378	21	AAAF8073	cDNA encoding huma
	43	16	2.6	401	21	AAAC03103	Human secreted pro
	44	16	2.6	422	21	AAZ80244	Human colon cancer
	45	16	2.6	484	21	AAAC01423	Human secreted pro

ALIGNMENTS

RESULT 1
AAF95006
ID AAF95006 standard; DNA; 606 BP.
XX
AC AAF95006;
DT 23-MAY-2001 (first entry)
XX
DE Human ovarian cancer associated coding sequence SEQ ID NO: 198.
XX
KW Human, ovarian cancer; vaccine; gene therapy; carcinoma; ds.
XX
OS Homo sapiens.
XX
FN WO200118046-A2.
XX
PD 15-MAR-2001.
XX
PF 08-SEP-2000; 2000WO-US24827.
XX
PR 10-SEP-1999; 99US-0394374.
PR 01-MAY-2000; 2000US-0561778.
PR 15-AUG-2000; 2000US-0640173.
PR 07-SEP-2000; 2000US-0656668.

(CORI-) CORIXA CORP.
Xu J, Stolk JA;
WPI; 2001-211395/21.
Isolated polypeptides associated with ovarian carcinomas, and the
nucleic acids that encode them, useful for the prevention diagnosis and
treatment of ovarian cancers -

RNA polymerase: in vitro transcriptional assay; probe; ss.

RNA polymerase; in vitro transcriptional assay; P	
Synthetic.	
Caenorhabditis elegans.	
WO9814574-A1.	
09-APR-1998.	
02-OCT-1997; 97WO-US17992.	
04-OCT-1996; 96US-0725459.	
{OKLA-} OKLAHOMA MEDICAL RES FOUND.	
Conaway JW, Conaway RC;	
WPI; 1998-286420/25.	
Isolated Elongin A and C and fragments - used for regulating transcriptional activation of RNA polymerase, particularly for use in in vitro assays and systems	
Example 7; Page 44; 185pp; English.	
The present sequence represents a probe used to isolate DNA encoding an Elongin A homologue of Caenorhabditis elegans. Elongin A stimulates the elongation activity of RNA polymerase II. Fragments of the rat Elongin A (AAW60772-76) have been found to each possess 50% or more of the Elongin A sequence activity exhibited by the wild type protein. The Elongin A sequence critical for binding to Elongins B and C is also found within these fragments. The Elongin polypeptides and fragments can be used for regulating the transcriptional activity of RNA polymerase. They can be used in in vitro transcriptional assays or systems.	
Sequence 51 BP; 13 A; 9 C; 17 G; 12 T; 0 other;	

Query Match 3.5%; Score 21; DB 19; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.27; Indels 0;
Matches 21; Conservative 0; Mismatches 0; Gaps 0;

QY 579 atctactgaagcctgttcttg 599
 |||||
Db 44 ATCTACTGAAGCCTGTTCTTG 24

RESULT	3
AAC76385/c	
ID	AAC76385 standard; cDNA; 2769 BP.
XX	
XX	AAC76385;
AC	
XX	
DT	08-FEB-2001 (first entry)
XX	
XX	

DE	Human	ORF A	ORF1540	polymerase
DE	Human	open reading frame	ORFX	detection; cytostatic; hepatotropic;
KW	vulnary;	antipsoriatic;	antiparkinsonian;	nootropic; neuroprotective;
KW	anticonvulsant;	osteopathic;	antiarrhythmic;	immunosuppressant; cardiatic;
KW	immunostimulant;	thrombolytic;	coagulant;	vasotropic; antidiabetic;
KW	hypotensive;	dermatological;	immunosuppressive;	antiinflammatory;
KW	antiviral;	antibacterial;	antifungal;	antirheumatic; antithyroid;
KW	antianaemic;	gene therapy;	cancer;	proliferative disorder; hypertension;
KW	cardiogenerative disorder;	osteoarthritis;	graft vs host disease;	
KW	cardiovascular disease;	diabetes mellitus;	hypothyroidism; SCID; AIDS;	
KW	cholesterol ester storage;	systemic lupus erythematosus;	infection;	
KW	severe combined immunodeficiency;	malaria;	autoimmune disorder; asthma;	
KW	allergy; aplastic anaemia;	nocturnal haemoglobinuria;	burn; wound;	
KW	bone damage;	cartilage damage;	antiinflammatory disease; coagulation;	
KW	thrombosis;	contraceptive;	ss.	
OS	Homio sapiens.			

Claim 18; Page 189; 189pp: English.

The present invention provides a number of coding sequences and proteins, the over-expression of which is associated with ovarian carcinoma/cancer. These can be used in the diagnosis, treatment and prevention of ovarian cancer, optionally by gene therapy or in the form of a vaccine. The present sequence is an example of one of these sequences.

Sequence 606 BP; 175 A; 147 C; 123 G; 161 T; 0 other;

Query Match	100.0%;	Score 606;	DB 22;	Length 606;
Best Local Similarity	100.0%;	Pred. No. 0;	Indels 0;	Gaps 0;

Malchines	000,	Consol	111	60
Qy	1	tgagttgcccccttacccccattccccagtgaaatttgcaattcctaagacgtgtttg		60

Db
1 tgagtttgcgccttaccoccatcccccagtcgaatatattgcaattcctaagaagcgtgttctt

QY
61
Db

121 tgctaaacatcttttaacgcacaggacagagcccccaaaagaataattatctatgccaa 180

181 atgtccataaacctgctgttgagaaaacctaccgcagatcttactgggcttcataagta 240

db 181 atgtccataaacactgctgttgagaaaacctaccgcgaggtactcttaacggtggtggttccatagggaa

241 agcttgcccttgcttgcttctctgtagatataaaaaataaagacactgcccagtcctcc 300
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Qy 301 ctcaacgtcccgagcaggctcaaggcaattccaataaacagtagaatgucataat
302 atcaactcccgaacaggcgcagaagcaattccaataaacagttagaatgaacactaa
303 atcaactcccgaacaggcgcagaagcaattccaataaacagttagaatgaacactaa

361 ttgatttcaaaatctcagcaactagaagaatgaccaaccatcctggttgccctgggactg 420
QY

[illegible]

Db 421 tctagtttttagcattgaaagtttcaggattccaggaaagccctcaggcctgggctgctgg 480

QY 481 tcacctagcagctgagggaactcttccataacagaattatgtcttcctctgcaatggagggag 540

Qy 541 tatactttaattgttaacatgtgaaaaacatctataaacatctactgaagcctgttcttgt 600

D_B
541 tatactttaatttgtaacatgtgaaacaactcctaataaacaccclccccgaagccgcgccc
501 ctggac g06
C:

Db 601 ctgcac 606

RESULT 2

ID AAV36272 standard; DNA; 51 BP.
XX
XX AAV36272.
AC

XX	08-SEP-1998	(first entry)
DT		
VV		

DE Probe used to isolate DNA encoding Elongin A homologues
XX Elongin A; rat; elongation activity; stimulation; RNA polymerase II;
vw probe used to isolate DNA encoding Elongin A homologues

KW binding; Elongin B; Elongin C; regulation, transcription

us-09-656-668-198.oli.rng

Wed Nov 7 09:21:18 2001

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XX FH Key Location/Qualifiers
XX CDS 33..818
XX FT /*tag= a
XX FT 33..92
XX FT /*tag= b
XX FT 93..818
XX FT /*tag= c
XX FT
XX FT
XX PN EP403114-A.
XX PD 19-DEC-1990.
XX PF 31-MAY-1990; 90EP-0305928.
XX PR 14-MAR-1990; 90US-0493588.
XX PR 15-JUN-1989; 89US-0366910.
XX PR 13-OCT-1989; 89US-0421201.
XX PR
XX PA (IMMU-) IMMUNEX CORP.
XX PI Park LS, Goodwin RG;
XX XX
XX DR WPI; 1990-377843/51.
XX DR P-PSDB; AAR08330.
XX XX
XX PT Mammalian interleukin-7 receptor DNA, protein and analogues -
XX PT used in therapy, diagnosis, assay and antibody production
XX PS Claim 4; Fig 3; 28pp; English.
XX XX
XX CC IL-7R gene product may be used in immunoassay or for affinity
XX CC purification eg. IL-7R, IL-1 or IL-1R. May also be useful in
XX CC regulation of immune response specifically of IL-7 in mammals
XX CC esp. humans.
XX XX
XX SQ Sequence 1427 BP; 438 A; 313 C; 313 G; 362 T; 1 other;

Query Match 3.0%; Score 18; DB 11; Length 1427;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 374 ctcagcaactagaagaat 391
| | | | | | | | | | | | | | | |
Dd 924 ctcagcaactagaagaat 941

RESULT 5
AAF21367
ID AAF21367 standard; DNA; 1658 BP.
XX AC
XX AAF21367;
XX XX
DT 14-MAR-2001 (first entry)
XX XX
XX DE Human low adenosine antisense oligonucleotide related sequence #2934.
XX DE
XX KW Low adenosine antisense oligonucleotide; phosphorothioate; allergy;
XX KW human; airway disorder; bronchoconstriction; lung inflammation;
XX KW surfactant depletion; respiratory; bronchodilator; antiinflammatory;
XX KW immunosuppressive; antiasthmatic; analgesic; hypotensive; cytostatic;
XX KW respiratory obstruction; pulmonary obstruction; impeded respiration;
XX KW surfactant hypoproduction; pulmonary vasoconstriction; asthma; RDS;
XX KW respiratory distress syndrome; pain; cystic fibrosis; allergic rhinitis;
XX KW pulmonary hypertension; emphysema; pulmonary transplantation rejection;
XX KW chronic obstructive pulmonary disease; pulmonary infection; bronchitis;
XX KW cancer; ss.
XX XX
XX OS Homo sapiens.
XX XX
XX PN WO200062736-A2.
XX XX

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XX PN WO200058473-A2.
XX XX
XX PD 05-OCT-2000.
XX XX
XX PF 31-MAR-2000; 2000WO-US08621.
XX XX
XX PR 31-MAR-1999; 99US-0127607.
XX PR 02-APR-1999; 99US-0127636.
XX PR 05-APR-1999; 99US-0127728.
XX PR 30-MAR-2000; 2000US-0540763.
XX XX
XX PA (CURA-) CURAGEN CORP.
XX XX
XX PI Shimkets RA, Leach M;
XX XX
XX DR WPI; 2000-602362/57.
XX DR P-PSDB; AAB421176.
XX XX
XX PT Novel nucleic acids and peptides derived from open reading frame X,
XX PT useful for treating e.g. cancers, proliferative disorders,
XX PT neurodegenerative disorders and cardiovascular disease -
XX XX
XX PS Claim 5; Page 3025-3027; 5507pp; English.
XX XX
XX CC AAC74446 to AAC77606 encode the proteins given in AAB40237 to AAB43397,
XX CC which represent the human ORFX open reading frames 1 to 3161. The ORFX
XX CC sequences have activities such as: cytostatic; hepatotropic; vulnarary;
XX CC antiproliferative; antiparkinsonian; nontropic; immunosuppressive;
XX CC osteopathic; anticonvulsant; antiarthritic; neuroprotective;
XX CC immunostimulant; cardiac; thrombotic; coagulant; vasotropic;
XX CC antidiabetic; hypotensive; dermatological; immunosuppressive;
XX CC antiinflammatory; antibacterial; antiviral; antifungal; antirheumatic;
XX CC antithyroid; and antianemic. The sequences can be used for determining
XX CC the presence of or predisposition to, or preventing or treating
XX CC pathological conditions associated with an ORFX-associated disorder. The
XX CC nucleic acids can be used to express ORFX proteins in gene therapy
XX CC vectors. The proteins and nucleic acids may be used to treat cancers,
XX CC proliferative disorders, neurodegenerative disorders, osteoarthritis,
XX CC graft vs host disease, cardiovascular disease, diabetes mellitus,
XX CC hypertension, hypothyroidism, cholesterol ester storage, systemic lupus
XX CC erythematosus, severe combined immunodeficiency (SCID), AIDS, viral,
XX CC bacterial or fungal infection, malaria, autoimmune disorders, asthma,
XX CC allergies, aplastic anaemia, burns, wounds, bone and cartilage damage,
XX CC nocturnal haemoglobinuria, antiinflammatory disease; to enhance
XX CC coagulation; to inhibit thrombosis; and as a contraceptive.
XX XX
XX SQ Sequence 2769 BP; 604 A; 789 C; 772 G; 603 T; 1 other;

Query Match 3.1%; Score 19; DB 21; Length 2769;
Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 425 agtttagcattgaaagt 443
| | | | | | | | | | | | | | | |
Dd 1419 AGTTTAGCATTGAAAGTT 1401

RESULT 4
AAQ06902
ID AAQ06902 standard; DNA; 1427 BP.
XX AC
XX AAQ06902;
XX XX
DT 04-MAR-1991 (first entry)
XX XX
XX DE Sequence encoding human IL-7 receptor clone H6.
XX XX
XX KW Interleukin-7 receptor; immune response; pre-B cell growth factor;
XX KW Lymphopietin; ds.
XX XX
XX OS Homo sapiens.

```

allergic disease; bronchoconstriction; inhibitor; antiinflammatory;
antiallergic; antiasthmatic; cycostatic; analgesic; impaired airway;
lung disease; ischaemic condition; pulmonary vasoconstriction; asthma;
respiratory distress syndrome; pain; cystic fibrosis; emphysema;
pulmonary hypertension; chronic obstructive pulmonary disease; COPD;
pulmonary diseases; metastasis; ss.

allergic diseases
antiallergic; a
lung disease; i
respiratory dis
pulmonary hyper
cancer; leukaemia
Homo sapiens.
W02000009525-A2.

24-FEB-2000. 99WO-US17712.
03-AUG-1999; 98US-0095212.
03-AUG-1998; 98US-0095212.
UNIV EAST CAROLINA.

Nyce JW;
DOI: 10.1002-205971/18.

New antisense oligonucleotides useful for treating e.g. pulmonary vasoconstriction, inflammation, allergies, asthma, hypertension, bronchitis, emphysema, respiratory distress syndrome, cancer.

Disclosure; Page 1284-1285; 1343pp; English.

The present invention describes a new composition comprising an antisense oligonucleotide (ON) with low adenosine (up to 15%), which targets nucleic acids involved in bronchoconstriction, allergies, and/or inflammation. The ON can have antiinflammatory, antiallergic, antiasthmatic, cytostatic and analgesic activities. The compositions are useful for the treatment of diseases associated with inflammation, impaired airways, including lung disease and diseases whose secondary effects affect the lungs of a subject. They can be used for treating e.g. ischaemic conditions, pulmonary vasoconstriction, allergies, asthma, impaired respiration, respiratory distress syndrome, pain, cystic fibrosis, pulmonary hypertension, emphysema, chronic obstructive pulmonary disease (COPD), and cancers such as leukaemias, lymphomas, carcinomas, and cancers which may metastasize to the lungs, including breast and prostate cancer. The reduction of the adenosine content of the ONs reduces side effects. The A-containing ONs break down with the release of deoxyadenosine which activates adenosine receptors causing the bronchoconstriction and inflammation. AAA32313 to AAA35312 represent nucleotide sequences given in the sequence listing from the present invention, which correspond to SEQ ID NO:1 to 2815, and then the sequences 185 sequences are also called SEQ ID NO:1 to 185, but the sequences differ from the previously named sequences. SEQ ID NO:11 to 1680 (AAA32323 to AAA33992) are specifically claimed ONs from the present invention. N.B. Sequences given in the disclosure of the present invention do not match up with their corresponding SEQ ID NO: sequences in the sequence listing.

Sequence 1658 BP; 493 A; 364 C; 363 G; 438 T; 0 other;

Length 1658; pp 21.

Query Match	3.0%;	Score 18;	DB 41;	Length 1000;
Best Local Similarity	100.0%;	Pred. No. 12;		
18. Conservative	0;	Mismatches	0;	Indels 0; Gaps 0;

374 ctcagcaactagaagaat 391
|||
1025

RESULT	7
AAQ06901	
ID	AAQ06901 standard; cdna; 1674 bp.
XX	
AC	AAQ06901;

26-OCT-2000.
24-MAR-2000; 2000WO-US08020.
06-APR-1999; 99US-0127958.
(DYEC-) UNIV EAST CAROLINA.
(NYCE/) NYCE J W.
Nyce JW;
WPI; 2000-679539/66.
Low adenosine (A) content antisense oligonucleotides which do not
trigger adenosine receptors during metabolism, useful e.g. for treating
cancers and respiratory obstructions -
Paco 1370. 1592pp. English.

The present invention describes low adenosine (A) content antisense oligonucleotides and compositions (I) comprising them. In the antisense oligonucleotides the A is replaced by a 'universal' or alternative base. (I) can have respiratory, bronchodilator, antiinflammatory, analgesic, immunosuppressive, antiasthmatic, hypotensive and cytostatic activities. The antisense oligonucleotides and (I) can be used to down-regulate the expression and/or activity of target polypeptides associated with lung/respiratory disorders and malignancies, such as stimulating and activating peptide factors and transmitters, transcription factors, immunoglobulins and antibodies, antibody receptors, cytokines and chemokines, endogenously produced specific and non-specific enzymes, binding proteins, adhesion molecules and their receptors, cytokine and chemokine receptors, adenosine receptors, bradykinin receptors, central nervous system (CNS) and peripheral nervous and non-nervous system receptors, CNS and peripheral nervous and non-nervous system peptide transmitters, defensins, growth factors, vasoactive peptides and receptors, binding proteins and malignancy associated proteins. The antisense oligonucleotides may be used in this way to treat disorders including respiratory obstruction (especially pulmonary obstruction and/or bronchoconstriction) and/or lung inflammation, allergy(ies) and/or bronchoconstriction) which are associated with a disease or condition selected from pulmonary vasoconstriction, inflammation, allergies, asthma, impaired respiration, respiratory distress syndrome (RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary hypertension, emphysema, chronic obstructive pulmonary disease (COPD), and/or cancer. AAF18434 to AAF21543 represent human polynucleotide fragments and antisense oligonucleotides used in the exemplification of the present invention.

CC the present environment
XX
XX

SQ Sequence 1658 BP; 493 A; 304 C; 303 G; 168 T
 Query Match 3.0%; Score 18; DB 21; Length 1658;
 Best Local Similarity 100.08; P. Num. 12;
 Mismatches 0; Indels 0; Gaps 0;

Qy 374 ctcagcaactagaagaat 391
|||||

RESULT 6
AAA35245
IN AAA35245 standard. DNA: 1658 BP.

XX AC AAA35245:

XX DT 28-JUL-2000 (first entry)

DI 20 OCT 2006 (continued)
XX
XX

DE	Human adenosine receptor related polynucleotide and
XX	low adenosine antisense oligonucleotide;
KW	Human; adenosine receptor; low adenosine receptor; inflammation; allergy;

us-09-656-668-198.oli.rng

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IP-10; CRG-2; CTLA-8; herpesvirus; Saimiri; ds.

```

XX KW IP-10; CRG-2; CTLA-8; herpesvirus; Saimiri; ds.
XX OS Homo sapiens.
XX Key Location/Qualifiers
XX FH 52..2037
XX FT CDS /*tag= a
XX FT
XX PN WO9707198-A2.
XX PN
XX XX 27-FEB-1997.
XX PD
XX PF 08-AUG-1996; 96WO-US12897.
XX PF
XX PR 08-AUG-1996; 96WO-US12897.
XX PR
XX PA (GEM ) GENETICS INST INC.
XX PA
XX PI Carlin M, Jacobs K, Kelleher K, McCoy JM;
XX PI
XX XX WPI; 1997-165283/15.
XX XX P-PSDB; AAW28510.
XX DR
XX XX Polynucleotide(s) encoding proteins for treating, preventing and
XX PT ameliorating medical conditions - obtained from human activated
XX PT peripheral blood mononuclear cell, and murine adult thymus libraries
XX PT
XX XX Claim 11; Page 39-42; 61pp; English.
XX PS
XX XX This sequence was isolated from a human activated PBMC library using
XX CC a trap selecting for nucleotides encoding secreted proteins, and
XX CC encodes a protein having homology with Drosophila leucine-rich
XX CC repeat proteins.
XX CC
XX XX Sequence 2582 BP; 726 A; 653 C; 508 G; 695 T; 0 other;
XX SQ

Query Match 3.0%; Score 18; DB 18; Length 2582;
Best Local Similarity 100.0%; Pred. No. 13;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 286 ctgccagtcctccctcctc 303
Db 2328 CTGCCAGTCCTCCCTC 2311

RESULT 9
AAV15587/c
ID AAV15587 standard; cDNA; 2635 BP.
XX AC AAV15587;
XX AC
XX XX 22-MAY-1998 (first entry)
XX DT
XX DE Human B-cell activation and survival antigen-1 cDNA.
XX DE
XX KW Human; B-cell activation and survival antigen-1; BAS-1; diagnosis;
XX KW treatment; abnormal B-cell response; abnormal proliferation;
XX KW cancerous condition; degenerative condition; autoimmune disease;
XX KW allergic response; ss.
XX KW
XX OS Homo sapiens.
XX OS
XX FH Key Location/Qualifiers
XX FT CDS 97..2082
XX FT /*tag= a
XX FT /product= BAS-1
XX FT
XX XX WO9744452-Al.
XX PN
XX PD 27-NOV-1997.
XX PD
XX XX 15-MAY-1997; 97WO-US07648.
XX PF

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XX DF 04-MAR-1991 (first entry)
XX DE
XX DE Sequence encoding human IL-7 receptor clone 1120.
XX KW Interleukin-7 receptor; immune response; pre-B cell growth factor;
XX KW Lymphopoietin; ds.
XX KW
XX OS Homo sapiens.
XX OS
XX FH Key Location/Qualifiers
XX FT CDS 23..1399
XX FT /*tag= a
XX FT sig_peptide 23..82
XX FT /*tag= b
XX FT mat_peptide 83..1399
XX FT /*tag= c
XX FT misc_RNA 740..814
XX FT /*tag= d
XX FT /label= Encodes transmembrane region
XX FT
XX PN EP403114-A.
XX PN
XX XX 19-DEC-1990.
XX XX
XX XX 31-MAY-1990; 90EP-0305928.
XX PF
XX PR 14-MAR-1990; 90US-0493588.
XX PR
XX PR 15-JUN-1989; 89US-0366910.
XX PR
XX PR 13-OCT-1989; 89US-0421201.
XX PR
XX PA (IMMUNEX) IMMUNEX CORP.
XX PA
XX PI Park LS, Goodwin RG;
XX PI
XX XX WPI; 1990-377843/51.
XX DR P-PSDB; AAR08329.
XX DR
XX XX Mammalian interleukin-7 receptor DNA, protein and analogues -
XX PT used in therapy, diagnosis, assay and antibody production
XX PT
XX PS Claim 4; Fig 2; 28pp; English.
XX PS
XX XX IL-7R gene product may be used in immunoassay or for affinity
XX CC purification eg. IL-7R, IL-1 or IL-1R. May also be useful in
XX CC regulation of immune response specifically of IL-7 in mammals
XX CC esp. humans.
XX CC
XX XX Sequence 1674 BP; 509 A; 366 C; 363 G; 436 T; 0 other;
XX SQ

Query Match 3.0%; Score 18; DB 11; Length 1674;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 374 ctacagcaactagaagaat 391
Db 1008 ctacagcaactagaagaat 1025

RESULT 8
AAT87428/c
ID AAT87428 standard; cDNA; 2582 BP.
XX AC
XX AC AAT87428;
XX AC
XX DT 29-DEC-1997 (first entry)
XX DT
XX XX Clone J422.
XX DE
XX XX J5; J422; L105; H174-10; H174-43; B18; cytokine; PBMC;
XX KW peripheral blood mononuclear cell; disintegrin; metallo-protein;
XX KW Drosophila; leucine-rich repeat; monocyte; chemoattractant;
XX KW

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XX PS Claim 2; Page 10-13; 15pp; Japanese.
XX PS The present sequence encodes a B cell surface protein. The protein
CC promotes growth of B cells as well as inhibition and induction of B
CC cell apoptosis. The protein and the DNA are useful in the treatment
CC and as a diagnostic agent for B-cell antibody production related
CC diseases e.g. autoimmune diseases, allergic diseases, asthma and
CC atopic dermatitis.
XX SQ Sequence 2697 BP; 757 A; 680 C; 533 G; 727 T; 0 other;

Query Match 3.0%; Score 18; DB 20; Length 2697;
Best Local Similarity 100.0%; Pred. No. 13;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 286 ctgccagtcctccctc 303
DB 2419 CTGCCAGTCCTCCCTC 2402
|||||

RESULT 11
AAV83816/C
ID AAV83816 standard; cDNA; 2697 BP.
XX AC AAV83816;
XX DT 03-MAR-1999 (first entry)
XX DE cDNA encoding a B cell surface protein.
XX KW B cell surface protein; growth; inhibition; induction; B cell apoptosis;
XX OS Homo sapiens.
XX PN AAV83816;
XX PD 2373 CTGCCAGTCCTCCCTC 2356
QY 286 ctgccagtcctccctc 303
DB 2373 CTGCCAGTCCTCCCTC 2356
|||||

RESULT 10
AAV83816/C
ID AAV83816 standard; cDNA; 2697 BP.
XX AC AAV83816;
XX DT 03-MAR-1999 (first entry)
XX DE cDNA encoding a B cell surface protein.
XX KW B cell surface protein; growth; inhibition; induction; B cell apoptosis;
XX OS Homo sapiens.
XX FH Key Location/Qualifiers
XX CDS 143..2128
XX FT /*tag= a
XX FT sig_peptide 143..203
XX FT /*tag= b
XX FT mat_peptide 204..2125
XX FT /*tag= c
XX JPI0313870-A.
XX PN
XX PD 02-DEC-1998.
XX PF 22-MAY-1997; 97JP-0132592.
XX PR 22-MAY-1997; 97JP-0132592.
XX PA (MITU ) MITSUBISHI CHEM CORP.
XX WPI; 1999-084647/08.
XX P-PSDB; AAW87556.
XX New DNA sequence encoding B cell surface protein - useful as
XX diagnostic agent and for treating auto-immune diseases, allergic
XX diseases, asthma and atopic dermatitis
PT

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SQ Sequence 2775 BP; 792 A; 694 C; 565 G; 724 T; 0 other;

Query Match 3.0%; Score 18; DB 18; Length 2775;

Best Local Similarity 100.0%; Pred. No. 13; 0; Indels 0; Gaps 0;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 286 ctgccagtcctccctc 303

DB 2417 CTGCCAGTCCCTCCCTC 2400

RESULT 12

AAA08399/c

ID AAA08399 standard; DNA; 2775 BP.

XX

AC AAA08399;

XX

XX

DT 12-JUL-2000 (first entry)

XX

DE Human RP105 nucleotide sequence SEQ ID NO:2.

XX

XX Human; RP105; agonist; antagonist; allergy; allergic disease;

KW identification; antiasthmatic; dermatological; antiinflammatory;

KW cytotatic; immunosuppressive; neuroprotective; antianaemic;

KW antidiabetic; asthma; atopic dermatitis; B cell neoplasm;

KW chronic lymphocyte leukaemia; hairy cell leukaemia; myeloma;

KW polymphocytic leukaemia; autoimmune disease; rheumatoid arthritis;

KW systemic lupus erythematosus; multiple sclerosis; diabetes;

KW acquired haemolytic anaemia; ds.

XX

OS Homo sapiens.

XX

XX WO200012130-A1.

PN

XX

XX 09-MAR-2000.

PD

XX

XX 24-AUG-1999; 99WO-US19336.

PF

XX

XX 27-AUG-1998; 98US-0098030.

PR

XX

XX (SMIK) SMITHKLINE BEECHAM CORP.

PA

XX

XX (SMIK) SMITHKLINE BEECHAM PLC.

PI

XX

XX Harrop JA, Holmes SD, Roshak AK;

XX

XX WPI; 2000-375617/32.

DR

XX

XX P-PSDB; AAY82527.

DR

XX

XX

New anti-murine B cell antigen RP105 antibody for treating or

preventing allergic disease such as asthma, B cell neoplasms, for

example leukemias, and auto immune diseases such as systemic lupus

erythematosus

Example; Page 26-28; 30pp; English.

The present invention describes an anti-murine B cell antigen RP105

antibody (I). The present sequence encodes human RP105, which is used

in the exemplification of the present invention. (I) has antiasthmatic,

dermatological, antiinflammatory, cytostatic, immunosuppressive,

neuroprotective, antianaemic and antidiabetic activities. (I) can be

used for in the treatment and prevention of allergic disease, including

asthma and atopic dermatitis, B cell neoplasms, including chronic

lymphocyte leukaemia, hairy cell leukaemia, polymphocytic leukaemia,

myelomas, autoimmune diseases such as systemic lupus erythematosus,

rheumatoid arthritis, multiple sclerosis, acquired haemolytic anaemia

and diabetes.

Sequence 2775 BP; 792 A; 694 C; 565 G; 724 T; 0 other;

Query Match 3.0%; Score 18; DB 21; Length 2775;

Best Local Similarity 100.0%; Pred. No. 13;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 286 ctgccagtcctccctc 303

DB 2417 CTGCCAGTCCCTCCCTC 2400

RESULT 13

AAF21368

ID AAF21368 standard; DNA; 3191 BP.

XX

AC AAF21368;

XX

DT 14-MAR-2001 (first entry)

XX

DE Human low adenosine antisense oligonucleotide related sequence #2935.

XX

XX Low adenosine antisense oligonucleotide; phosphorothioate; allergy;

KW human; airway disorder; bronchoconstriction; lung inflammation;

KW surfactant depletion; respiratory; bronchodilator; antiinflammatory;

KW immunosuppressive; antiasthmatic; analgesic; hypotensive; cytostatic;

KW respiratory obstruction; pulmonary obstruction; impeded respiration;

KW surfactant hypoproduction; pulmonary vasoconstriction; asthma; RDS;

KW respiratory distress syndrome; pain; cystic fibrosis; allergic rhinitis;

KW pulmonary hypertension; emphysema; pulmonary transplantation rejection;

KW chronic obstructive pulmonary disease; pulmonary infection; bronchitis;

KW cancer; ss.

XX

OS Homo sapiens.

XX

XX WO200062736-A2.

PN

XX

XX 26-OCT-2000.

PD

XX

XX 24-MAR-2000; 2000WO-US08020.

PF

XX

XX 06-APR-1999; 99US-0127958.

PR

XX

XX (UYEC-) UNIV EAST CAROLINA.

PA

XX

XX (NYCE/) NYCE J W.

PI

XX

XX Nyce JW;

XX

XX WPI; 2000-679539/66.

DR

XX

XX Low adenosine (A) content antisense oligonucleotides which do not

trigger adenosine receptors during metabolism, useful e.g. for treating

cancers and respiratory obstructions

PT

XX

XX Disclosure; Page 1370-1371; 1592pp; English.

PS

The present invention describes low adenosine (A) content antisense

oligonucleotides and compositions (I) comprising them. In the antisense

oligonucleotides the A is replaced by a 'Universal' or alternative base.

(I) can have respiratory, bronchodilator, antiinflammatory, analgesic,

immunosuppressive, antiasthmatic, hypotensive and cytostatic activities.

The antisense oligonucleotides and (I) can be used to down-regulate the

expression and or activity of target polypeptides associated with

lung/respiratory disorders and malignancies, such as stimulating and

activating peptide factors and transmitters, transcription factors,

immunoglobulins and antibodies, antibody receptors, cytokines and

chemokines, endogenously produced specific and non-specific enzymes,

binding proteins, adhesion molecules and their receptors, cytokine and

chemokine receptors, adenosine receptors, bradykinin receptors, central

nervous system (CNS) and peripheral nervous and non-nervous system

receptors, CNS and peripheral nervous and non-nervous system peptide

transmitters, defensins, growth factors, vasoactive peptides and

receptors, binding proteins and malignancy associated proteins. The

antisense oligonucleotides may be used in this way to treat disorders

including respiratory obstruction (especially pulmonary obstruction

and/or bronchoconstriction) and/or lung inflammation, allergy(ies)

and/or surfactant hypoproduction which are associated with a disease or

condition selected from pulmonary vasoconstriction, inflammation,

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CC allergies, asthma, impeded respiration, respiratory distress syndrome
 CC (RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary
 CC hypertension, emphysema, chronic obstructive pulmonary disease (COPD),
 CC pulmonary transplantation rejection, pulmonary infections, bronchitis,
 CC and/or cancer. AAF18434 to AAF21543 represent human polynucleotide
 CC fragments and antisense oligonucleotides used in the exemplification of
 CC the present invention.
 XX Sequence 3191 BP; 947 A; 674 C; 663 G; 907 T; 0 other;
 SQ

Query Match 3.0%; Score 18; DB 21; Length 3191;
 Best Local Similarity 100.0%; Pred. No. 13;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 374 ctcagcaactagaagaat 391
 Db 2541 ctcagcaactagaagaat 2558
 |||||

RESULT 14
 AAA35246
 ID AAA35246 standard; DNA; 3191 BP.
 XX
 AC AAA35246;
 DT - 28-JUL-2000 (first entry)
 XX Human adenosine receptor related polynucleotide 2nd SEQ ID NO:120.
 DE
 KW Human; adenosine receptor; low adenosine antisense oligonucleotide;
 KW phosphorothioate; impaired respiration; inflammation; allergy;
 KW allergic disease; bronchoconstriction; inhibitor; antiinflammatory;
 KW antiallergic; antiasthmatic; cytosolic; analgesic; impaired airway;
 KW lung disease; ischaemic condition; pulmonary vasoconstriction; asthma;
 KW respiratory distress syndrome; pain; cystic fibrosis; emphysema;
 KW pulmonary hypertension; chronic obstructive pulmonary disease; COPD;
 KW cancer; leukaemia; lymphoma; carcinoma; metastasis; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200009525-A2.
 XX
 PD 24-FEB-2000.
 XX
 PF 03-AUG-1999; 99WO-US17712.
 XX
 PR 03-AUG-1998; 98US-0095212.
 XX
 PA (UYEC-) UNIV EAST CAROLINA.
 XX
 PI Nyce JW;
 XX
 PP WPI: 2000-205971/18.
 XX
 DR New antisense oligonucleotides useful for treating e.g. pulmonary
 PT vasoconstriction, inflammation, allergies, asthma, hypertension,
 PT bronchitis, emphysema, respiratory distress syndrome, ischemia or
 PT cancers.
 XX
 PS Disclosure: Page 1285-1286; 1343pp; English.
 XX
 CC The present invention describes a new composition comprising an
 CC antisense oligonucleotide (ON) with low adenosine (up to 15%), which
 CC targets nucleic acids involved in bronchoconstriction, allergies, and/or
 CC inflammation. The ON can have antiinflammatory, antiallergic,
 CC antiasthmatic, cytosolic and analgesic activities. The compositions are
 CC useful for the treatment of diseases associated with inflammation,
 CC impaired airways, including lung disease and diseases whose secondary
 CC effects afflict the lungs of a subject. They can be used for treating
 CC e.g. ischaemic conditions, pulmonary vasoconstriction, allergies,
 CC asthma, impeded respiration, respiratory distress syndrome, pain, cystic
 CC fibrosis, pulmonary hypertension, emphysema, chronic obstructive

CC pulmonary disease (COPD), and cancers such as leukaemias, lymphomas,
 CC carcinomas, and cancers which may metastasise to the lungs, including
 CC breast and prostate cancer. The reduction of the adenosine content of
 CC the ONs reduces side effects. The A-containing adenosine receptors causing the
 CC release of deoxyadenosine which activates adenosine receptors causing the
 CC bronchoconstriction and inflammation. AAA32313 to AAA35312 represent the
 CC nucleotide sequences given in the sequence listing from the present
 CC invention, which correspond to SEQ ID NO:1 to 2815, and then the last
 CC 185 sequences are also called SEQ ID NO:1 to 185, but the sequences
 CC differ from the previously named sequences. SEQ ID NO:11 to 1680
 CC (AAA32323 to AAA33992) are specifically claimed ONs from the present
 CC invention. N.B. Sequences given in the disclosure of the present
 CC invention do not match up with their corresponding SEQ ID NO: sequences
 CC given in the sequence listing.
 XX Sequence 3191 BP; 947 A; 674 C; 663 G; 907 T; 0 other;
 SQ

Query Match 3.0%; Score 18; DB 21; Length 3191;
 Best Local Similarity 100.0%; Pred. No. 13;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 374 ctcagcaactagaagaat 391
 Db 2541 ctcagcaactagaagaat 2558
 |||||

RESULT 15
 AAT42063
 ID AAT42063 standard; DNA; 1830121 BP.
 XX
 AC AAT42063;
 DT 14-SEP-1999 (first entry)
 XX Haemophilus influenzae complete genome sequence.
 DE
 KW Genome; bacterium; Haemophilus influenzae; computer readable medium;
 KW expression modulating fragment; regulation; gene expression; vector;
 KW organism; open reading frame; ORF; ds.
 XX
 OS Haemophilus influenzae.
 XX
 PN WO9633276-A1.
 XX
 PD 24-OCT-1996.
 XX
 PF 22-APR-1996; 96WO-US05320.
 XX
 PR 07-JUN-1995; 95US-0487429.
 PR 21-APR-1995; 95US-0426787.
 PR 07-JUN-1995; 95US-0476102.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 PA (UYJO) UNIV JOHNS HOPKINS.
 XX
 PP Adams MD, Fleischmann RD, Smith HO, Venter JC, White O;
 WPI: 1996-485782/48.
 XX
 DR Haemophilus influenzae Rd genome recorded on computer readable
 PT medium - useful for identifying commercially important nucleic acid
 PT fragments by homology searching
 XX
 PS Claim 1; Page 77.2-77.1091; 1291pp; English.
 CC This sequence represents the complete genome sequence of the bacterium
 CC Haemophilus influenzae strain Rd. The invention relates to a computer
 CC readable medium (CRM) having recorded upon it the complete H. influenzae
 CC nucleotide sequence (1), a representative fragment of (1) or a nucleotide
 CC sequence at least 99% identical to (1). By providing the full-length
 CC genomic sequence in a computer readable form, it is possible to identify
 CC commercially important nucleic acid fragments and expression modulating

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CC fragments (EMFs) of the Haemophilus genome. The EMFs can be used to
 CC regulate the expression of a nucleic acid molecule. Vectors and altered
 CC organisms comprising the predicted ORFs can be used to produce any of the
 CC polypeptide fragments of the H. influenzae Rd genome.

XX
 SQ Sequence 1830121 BP; 567399 A; 350615 C; 347389 G; 564036 T; 682 other;

Query Match 3.0%; Score 18; DB 17; Length 1830121;
 Best Local Similarity 100.0%; Pred. No. 17;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 580 tctactgaagcctgttct 597
 |||||
 Db 408370 tctactgaagcctgttct 408387

Search completed: November 5, 2001, 23:48:56
 Job time: 2896 sec

GenCore version 4.5
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ow nucleic - nucleic search. using sw model

Run on: November 5, 2001, 22:16:40 ; Search time 87.77 Seconds
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Title: US-09-656-668-198
Perfect score: 606
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Scoring table: IDENTITY_NUC
Gapox 10.0 Gapext 1.0

Searched: 351203 seqs, 113238999 residues
total number of hits satisfying chosen parameters: 702406

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Minimum DB seq length: 0
Maximum DB seq length: 20000000000
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query			DB	ID	Description
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1	82.4	13.6	80246	4	US-09-078-294-4	Sequence 4, Appli
2	82.4	13.6	80595	4	US-09-078-294-3	Sequence 3, Appli
c 3	64.8	10.7	7505	4	US-09-078-294-13	Sequence 13, Appli
c 4	60.4	10.0	6314	1	US-08-211-430-1	Sequence 1, Appli
5	46.8	7.7	279	3	US-09-157-177-120	Sequence 120, App
6	41	6.8	1750	4	US-09-276-531-34	Sequence 34, Appli
c 7	38.6	6.4	11288	4	US-08-646-301A-1	Sequence 1, Appli
c 8	38.6	6.4	11288	4	US-08-481-968A-4	Sequence 4, Appli
c 9	36.6	6.0	3176	2	US-08-910-733-17	Sequence 17, Appli
c 10	36.6	6.0	3176	2	US-08-910-884-17	Sequence 17, Appli
11	35.6	5.9	11531-1	1	US-08-068-945A-1	Sequence 1, Appli
12	35.6	5.9	11531	1	US-08-442-806-1	Sequence 3, Appli
13	35.4	5.8	1481	3	US-09-429-323-3	Sequence 31, Appli
14	35.4	5.8	3073	2	US-07-688-352C-31	Sequence 31, Appli
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16	35.4	5.8	3073	3	US-09-146-249A-31	Sequence 31, Appli
17	35.4	5.8	3073	3	US-08-206-188B-31	Sequence 31, Appli
18	35.4	5.8	3073	5	PCR-US91-02714-30	Sequence 20, Appli
19	34.4	5.7	246240	2	US-08-724-394A-20	Sequence 20, Appli
20	34.4	5.7	246240	2	US-08-724-394A-21	Sequence 21, Appli
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c 22	34	5.6	2218	2	US-08-421-044-1	Sequence 1, Appli
c 23	32.8	5.4	8353	3	US-08-611-587-1	Sequence 1, Appli
c 24	31.8	5.2	1524	4	US-08-840-767-3	Sequence 3, Appli
c 25	31.2	5.1	5427	3	US-09-009-913-2	Sequence 2, Appli
c 26	31.2	5.1	5510	3	US-09-009-913-3	Sequence 3, Appli
c 27	31.2	5.1	5567	3	US-09-009-913-4	Sequence 4, Appli

Sequence 17, Appl
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Sequence 3, Appl
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Sequence 3, Appl
Sequence 5, Appl
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Sequence 5, Appl
Sequence 4, Appl
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Sequence 1, Appl
Sequence 97, Appl
Sequence 3, Appl
Sequence 5, Appl
Sequence 2, Appl

ALIGNMENTS

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1
RESULT
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; Sequence 4, Application US/09078294
; Patent No. 6265211
; GENERAL INFORMATION:
; APPLICANT: Choo, Kong-Hong Andy
; APPLICANT: Du Sart, Desiree
; APPLICANT: Cancilla, Michael R.
; TITLE OF INVENTION: A NOVEL NUCLEIC ACID MOLECULE
; FILE REFERENCE: Davies Col
; CURRENT APPLICATION NUMBER: US/09/078,294
; CURRENT FILING DATE: 1998-05-13
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 4
; LENGTH: 80246
; TYPE: DNA
; ORGANISM: Nucleotide sequence of NC-contig
US-09-078-294-4

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[illegible]

RESULT 2
US-09-078-294-3
; Sequence 3, Application US/09078294
; Patent No. 6265211
; GENERAL INFORMATION:
; APPLICANT: Choo, Kong-Hong Andy
; APPLICANT: Du Sart, Desiree
; APPLICANT: Cancilla, Michael R.

```

; TITLE OF INVENTION: A NOVEL NUCLEIC ACID MOLECULE
; FILE REFERENCE: Davies Col
; CURRENT APPLICATION NUMBER: US/09/078,294
; CURRENT FILING DATE: 1998-05-13
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
; LENGTH: 80595
; TYPE: DNA
; ORGANISM: Nucleotide sequence of HC-contig
US-09-078-294-3

Query Match      13.6%; Score 82.4; DB 4; Length 80595;
Best Local Similarity 63.8%; Pred. No. 4.2e-17;
Matches 125; Conservative 0; Mismatches 71; Indels 0; Gaps 0;

QY 72 tgggtgggaacatgctactggtcattgcatgcatgcatagagggcagtaagtgtctgtaaacatc 131
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DB 66788 tgaatgggtgggtggtactggcattgctgtagtgggtggagaccagagatgctgttaaacatc 66847

QY 132 ttccaagcacaggagacagcccccacaaaagagaattatctagcccaaatgtccataac 191
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 66848 ccgcgaagcacaggacagtcctccgcacacaagaattatctgcccccaaatcatcagt 66907

QY 192 actgctgttgagaaaacctccgcaggatcttactgggtcttcataaggcttgccttt 251
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB -66908 gccaaagtgtgagaacctcattctgactgtctcttccctcttcactgtcttaatacaactgtt 66967

QY 252 gttctggcttcgttag 267
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DB 66968 gttcttcagcattag 66983

RESULT 3
US-09-078-294-13/c
; Sequence 13, Application US/09078294
; Patent No. 6265211
; GENERAL INFORMATION:
; APPLICANT: Choo, Kong-Hong Andy
; APPLICANT: Du Sart, Desiree
; APPLICANT: Cancilla, Michael R.
; TITLE OF INVENTION: A NOVEL NUCLEIC ACID MOLECULE
; FILE REFERENCE: Davies Col
; CURRENT APPLICATION NUMBER: US/09/078,294
; CURRENT FILING DATE: 1998-05-13
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 13
; LENGTH: 7505
; TYPE: DNA
; ORGANISM: BAC-F2 contig 9
US-09-078-294-13

Query Match      10.7%; Score 64.8; DB 4; Length 7505;
Best Local Similarity 72.1%; Pred. No. 1e-11;
Matches 98; Conservative 0; Mismatches 37; Indels 1; Gaps 1;

QY 76 tggggaacatgctactggtcattgcatgcatagagggcagtaagtgtgtaaacatctttc 135
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DB 4460 TTGGGAATTTCGTACAGACATCTAATAATTAGAGCGTAGGGATATTGTTAAATGTCCTAC 4401

QY 136 aacgcacaggacagagccccacaaaagagaattatctagcccccaaatgtccataaacctg 195
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 4400 ACTGCACATGACAGACCCACATAAACAGAGAATTGTATAGCCCCAAAATGT-CATTAGTGTG 4342

QY 196 ctgttgagaaaccta 211
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DB 4341 ACGTTTGAGAAAAATATA 4326

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APPLICANT: Nilsson, Jeanette
 APPLICANT: Tornell, Jan
 TITLE OF INVENTION: Genomic DNA Sequences
 TITLE OF INVENTION: Encoding Human BSSL/CBL
 NUMBER OF SEQUENCES: 58
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: White & Case
 STREET: 1155 Avenue of the Americas
 CITY: New York
 STATE: New York
 COUNTRY: United States
 ZIP: 10036-2787
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patent Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/442,806
 FILING DATE:
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/068,945
 FILING DATE: 27-MAY-1993
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: SE 9201809-2
 FILING DATE: 11-JUN-1992
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: SE 9201826-6
 FILING DATE: 12-JUN-1992
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: SE 9202088-2
 FILING DATE: 03-JUL-1992
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: SE 9300902-5
 FILING DATE: 19-MAR-1993
 ATTORNEY/AGENT INFORMATION:
 NAME: Sterner, Richard J.
 REGISTRATION NUMBER: 35,372
 REFERENCE/DOCKET NUMBER: 1103326-052
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212)819-8783
 TELEFAX: (212)354-8113
 INFORMATION FOR SEQ ID NOS: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 11531 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: double
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 ORIGINAL SOURCE:
 ORGANISM: Homo sapiens
 TISSUE TYPE: Mammary gland
 FEATURE:
 NAME/KEY: CDS
 LOCATION: join(1653..1727, 4071..4221, 4307..4429, 4707
 LOCATION: ..4904, 6193..6323, 6501..6608, 6751..6868, 8335
 LOCATION: ..8521, 8719..8922, 10124..10321, 10650..11394)
 FEATURE:
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 LOCATION: join(1722..1727, 4071..4221, 4307..4429, 4707
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 LOCATION: ..8521, 8719..8922, 10124..10321, 10650..11391)
 OTHER INFORMATION: /EC_number= 3.1.1.1
 OTHER INFORMATION: /product= "Bile Salt-Stimulated Lipase"
 FEATURE:
 NAME/KEY: 5'UTR
 LOCATION: 1..1640
 FEATURE:
 NAME/KEY: TATA_signal
 LOCATION: 1611..1617
 FEATURE:

us-09-656-668-198.rni

Wed Nov 7 09:21:20 2001

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; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 3..1109
; US-08-474-379C-31

Query Match      5.8%; Score 35.4; DB 2; Length 3073;
Best Local Similarity 47.9%; Pred. No. 0.064;
Matches 102; Conservative 0; Mismatches 111; Indels 0; Gaps 0;

QY 70 cctgggtggggaacatgctactgcatcctaataatgcataagggcagtaatgctgctaaaca 129
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Db 280 CCGCGCGGGACAAAGAGTCAATCGCAACGACGAGGTTTGGGAACAGTAAATGGTTCATG 339

QY 130 tctttcaacgcacagacagagcccccacacaaagagaattatctagcccccaaatgtccata 189
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 340 TAAGCAACGGATATGTTTTCATCAACAGGAATGACACCAAGGAGATGTATTGTACACC 399

QY 190 acaactgctgttagaaaacctaccgcagagatcttactgggcttcattaggttaagcttgcc 249
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 400 AGACTGCCATAAAGAGAAATACCCAGGAGTACCTTCGCAGTGTAGGAGATGGAGAGA 459

QY 250 ttgttcggctctctgtagatatataaaataaag 282
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Db 460 CTGTGGAGTTTGATGTTGTTGAAGGAGAAAAGG 492
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Search completed: November 5, 2001, 22:58:05
Job time: 2485 sec